

## P. NT COOPERATION TREA

PCT

## NOTIFICATION OF ELECTION

(PCT Rule 61.2)

From the INTERNATIONAL BUREAU

To:

Assistant Commissioner for Patents  
 United States Patent and Trademark  
 Office  
 Box PCT  
 Washington, D.C.20231  
 ÉTATS-UNIS D'AMÉRIQUE

in its capacity as elected Office

|   |   |
|---|---|
| <b>Date of mailing (day/month/year)</b><br>02 November 1999 (02.11.99)        | <b>Applicant's or agent's file reference</b><br>DUNW/20631PC      |
| <b>International application No.</b><br>PCT/GB99/00963                        | <b>Priority date (day/month/year)</b><br>26 March 1998 (26.03.98) |
| <b>International filing date (day/month/year)</b><br>26 March 1999 (26.03.99) |   |
| <b>Applicant</b><br>WATTS, Colin  |   |

1. The designated Office is hereby notified of its election made:

☒ in the demand filed with the International Preliminary Examining Authority on:

14 October 1999 (14.10.99)

☐ in a notice effecting later election filed with the International Bureau on:2. The election ☒ was☐ was not

made before the expiration of 19 months from the priority date or, where Rule 32 applies, within the time limit under Rule 32.2(b).

The International Bureau of WIPO  
 34, chemin des Colombettes  
 1211 Geneva 20, Switzerland

Authorized officer

Juan Cruz

Facsimile No.: (41-22) 740.14.35

Telephone No.: (41-22) 338.83.38

# PATENT COOPERATION TREATY

# PCT

## INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

|  |   |   |
|--|---|---|
| Applicant's or agent's file reference<br><b>DUNW/20631PC</b> | <div style="display: flex; justify-content: space-between;"> <div style="text-align: center;"> <b>FOR FURTHER ACTION</b> </div> <div style="font-size: small;">             see Notification of Transmittal of International Search Report (Form PCT/ISA/220) as well as, where applicable, item 5 below.           </div> </div> |   |
| International application No.<br><b>PCT/GB 99/ 00963</b>     | International filing date (day/month/year)<br><div style="text-align: center;"><b>26/03/1999</b></div>  | (Earliest) Priority Date (day/month/year)<br><div style="text-align: center;"><b>26/03/1998</b></div> |
| Applicant<br><br><b>UNIVERSITY OF DUNDEE et al.</b>          |   |   |

This International Search Report has been prepared by this International Searching Authority and is transmitted to the applicant according to Article 18. A copy is being transmitted to the International Bureau.

This International Search Report consists of a total of 4 sheets.

☒ It is also accompanied by a copy of each prior art document cited in this report.

**1. Basis of the report**

a. With regard to the **language**, the international search was carried out on the basis of the international application in the language in which it was filed, unless otherwise indicated under this item.

☐ the international search was carried out on the basis of a translation of the international application furnished to this Authority (Rule 23.1(b)).

b. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international search was carried out on the basis of the sequence listing :

☐ contained in the international application in written form.

☐ filed together with the international application in computer readable form.

☒ furnished subsequently to this Authority in written form.

☒ furnished subsequently to this Authority in computer readable form.

☒ the statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.

☒ the statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished

2. ☒ **Certain claims were found unsearchable** (See Box I).

3. ☐ **Unity of invention is lacking** (see Box II).

4. With regard to the **title**,

☐ the text is approved as submitted by the applicant.

☒ the text has been established by this Authority to read as follows:

**USE OF INHIBITORS OF MAMMALIAN ASPARAGINYL ENDOPEPTIDASE FOR THERAPY OF AUTOIMMUNE DISEASE**

5. With regard to the **abstract**,

☒ the text is approved as submitted by the applicant.

☐ the text has been established, according to Rule 38.2(b), by this Authority as it appears in Box III. The applicant may, within one month from the date of mailing of this international search report, submit comments to this Authority.

6. The figure of the **drawings** to be published with the abstract is Figure No.

☒ as suggested by the applicant.

☐ because the applicant failed to suggest a figure.

☐ because this figure better characterizes the invention.

1

☐ None of the figures.

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/GB 99/00963

**Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)**

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 1-21  
because they relate to subject matter not required to be searched by this Authority, namely:  
Remark: Although claims 1-21 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. ☐ Claims Nos.:  
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

**Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)**

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

**Remark on Protest**

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

## INTERNATIONAL SEARCH REPORT

International Application No.

/GB 99/00963

## A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C07K5/06 C07K5/08 A61K38/55 C07K14/705 G01N33/68  
A01K67/027

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C07K A61K G01N A01K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

| Category ° | Citation of document, with indication, where appropriate, of the relevant passages   | Relevant to claim No. |
|------------|--|-----------------------|
| X          | CHEMICAL ABSTRACTS, vol. 128, no. 25,<br>22 June 1998 (1998-06-22)<br>Columbus, Ohio, US;<br>abstract no. 303708,<br>W C BENJAMIN ET AL.: "Caspase activation<br>in MCF7 cells responding to etoposide<br>treatment"<br>XP002113051<br>& MOL. PHARMACOL.,<br>vol. 53, no. 3, 1998, pages 446-450,<br>abstract<br><br>---<br>-/-- | 52, 53                |



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

° Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&amp;" document member of the same patent family

Date of the actual completion of the international search

23 August 1999

Date of mailing of the international search report

03/09/1999

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2

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Authorized officer

Masturzo, P

## INTERNATIONAL SEARCH REPORT

International Application No

GB 99/00963

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

| Category ° | Citation of document, with indication, where appropriate, of the relevant passages  | Relevant to claim No. |
|------------|---|-----------------------|
| X          | K K SHARMA ET AL.: "identification of new lens protease(s) using substrates having in vivo cleavage sites"<br>BIOCHEMICAL AND BIOPHYSICAL RESEARCH COMMUNICATIONS.,<br>vol. 218, 1996, pages 365-370, XP002113049<br>ACADEMIC PRESS INC. ORLANDO, FL., US<br>ISSN: 0006-291X<br>the whole document<br>---   | 55                    |
| A          | J M CHEN ET AL.: "Cloning, isolation and characterization of mammalian legumain, an asparaginyl endopeptidase"<br>JOURNAL OF BIOLOGICAL CHEMISTRY.,<br>vol. 272, no. 12,<br>21 March 1997 (1997-03-21), pages<br>80090-8098, XP002065429<br>AMERICAN SOCIETY OF BIOLOGICAL CHEMISTS,<br>BALTIMORE, MD., US<br>ISSN: 0021-9258<br>cited in the application<br>the whole document<br>---      | 1-56                  |
| P,X        | B MANOURY T AL.: "An asparaginyl endopeptidase processes a microbial antigen for class II MHC presentation"<br>NATURE.,<br>vol. 396, 17 December 1998 (1998-12-17),<br>pages 695-699, XP002113050<br>MACMILLAN JOURNALS LTD. LONDON., GB<br>ISSN: 0028-0836<br>the whole document<br>---  | 1-56                  |
| A          | CHEMICAL ABSTRACTS, vol. 119, no. 13,<br>27 September 1993 (1993-09-27)<br>Columbus, Ohio, US;<br>abstract no. 133992,<br>A A KEMBHAVI ET AL.: "The two cysteine endopeptidases of legume seeds; purification and characterization by use of specific fluorometric assays"<br>XP002113052<br>& ARCH. BIOCHEM. BIOPHYS.,<br>vol. 303, no. 2, 1993, pages 208-213,<br>abstract<br>---<br>-/-- | 43-45                 |

## INTERNATIONAL SEARCH REPORT

International Application No

/GB 99/00963

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT



| Category * | Citation of document, with indication, where appropriate, of the relevant passages   | Relevant to claim No. |
|------------|--|-----------------------|
| X          | <p>CHEMICAL ABSTRACTS, vol. 121, no. 17,<br/>24 October 1994 (1994-10-24)<br/>Columbus, Ohio, US;<br/>abstract no. 202741,<br/>M J ANDROLEWICZ ET AL.: "Human<br/>transporters associated with antigen<br/>processing possess a promiscuous binding<br/>site"<br/>XP002113053<br/>&amp; IMMUNITY,<br/>vol. 1, no. 1, 1994, pages 7-14,<br/>abstract</p> <p>-----</p> | 55                    |

## PATENT COOPERATION TREATY

## PCT

## INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

|  |  |   |
|--|--|---|
| Applicant's or agent's file reference<br><b>DUNW/P20631PC</b>  | <b>FOR FURTHER ACTION</b> See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)  |   |
| International application No.<br><b>PCT/GB99/00963</b>   | International filing date (day/month/year)<br><b>26/03/1999</b>  | Priority date (day/month/year)<br><b>26/03/1998</b> |
| International Patent Classification (IPC) or national classification and IPC<br><b>C07K5/06</b>  |  |   |
| Applicant<br><b>UNIVERSITY OF DUNDEE et al.</b>  |  |   |
| <p>1. This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.</p> <p>2. This REPORT consists of a total of 5 sheets, including this cover sheet.</p> <p><input checked="" type="checkbox"/> This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).</p> <p>These annexes consist of a total of 1 sheets.</p>   |  |   |
| <p>3. This report contains indications relating to the following items:</p> <ul style="list-style-type: none"><li>I <input checked="" type="checkbox"/> Basis of the report</li><li>II <input type="checkbox"/> Priority</li><li>III <input type="checkbox"/> Non-establishment of opinion with regard to novelty, inventive step and industrial applicability</li><li>IV <input type="checkbox"/> Lack of unity of invention</li><li>V <input checked="" type="checkbox"/> Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement</li><li>VI <input type="checkbox"/> Certain documents cited</li><li>VII <input type="checkbox"/> Certain defects in the international application</li><li>VIII <input checked="" type="checkbox"/> Certain observations on the international application</li></ul> |  |   |
| Date of submission of the demand<br><br><b>14/10/1999</b>  | Date of completion of this report<br><br><b>18. 07. 00</b>   |   |
| Name and mailing address of the international preliminary examining authority:<br><br>European Patent Office - P.B. 5818 Patentlaan 2<br>NL-2280 HV Rijswijk - Pays Bas<br>Tel. +31 70 340 - 2040 Tx: 31 651 epo nl<br>Fax: +31 70 340 - 3016   | Authorized officer<br><br><b>Masturzo, P</b><br><br>Telephone No. +31 70 340 2275<br> |   |

**INTERNATIONAL PRELIMINARY  
EXAMINATION REPORT**

International application No. PCT/GB99/00963

**I. Basis of the report**

1. This report has been drawn on the basis of (*substitute sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to the report since they do not contain amendments.*):

**Description, pages:**

1-72 as originally filed

**Claims, No.:**

1-51,52 (part) as originally filed

52 (part),53-56 as received on 23/03/2000 with letter of 22/03/2000

**Drawings, sheets:**

1/24-24/24 as originally filed

2. The amendments have resulted in the cancellation of:

- ☐ the description, pages:  
☐ the claims, Nos.:  
☐ the drawings, sheets:

3. ☐ This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)):

4. Additional observations, if necessary:



**INTERNATIONAL PRELIMINARY  
EXAMINATION REPORT**

International application No. PCT/GB99/00963

**V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement**

**1. Statement**

|                               |      |        |       |
|-------------------------------|------|--------|-------|
| Novelty (N)                   | Yes: | Claims | 1-56  |
|                               | No:  | Claims |       |
| Inventive step (IS)           | Yes: | Claims | 1-56  |
|                               | No:  | Claims |       |
| Industrial applicability (IA) | Yes: | Claims | 22-56 |
|                               | No:  | Claims | 1-21  |

**2. Citations and explanations**

**see separate sheet**

**VIII. Certain observations on the international application**

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:

**see separate sheet**

**INTERNATIONAL PRELIMINARY  
EXAMINATION REPORT - SEPARATE SHEET**

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International application No. PCT/GB99/00963

Reference is made to the following document/s/:

- D1: CA 128:303708 (March 1998);
- D2: Bioch. Biophys. Res. Comm. 218, 365-370 (1996);
- D3: CA 121: 202741 (1994).

**Re Item V**

**Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement**

1) D1 (see text) discloses compounds similar to those included in the formula of claim 52 and with inhibitory activity against caspase, a known thiol protease. No reference to AEP inhibitory activity is made. However, these compounds do not exhibit an asparagine residue prior to the CH<sub>2</sub>F residue which is the proper inhibiting group. The compound containing an asparagine described in the abstract (taken over from the abstract in the complete publication) owes its existence to a printing error.

The suggestion to use an asparagine residue in lieu of an aspartic acid residue is not present in the prior art either. Therefore claims 52 and 53 are new and inventive under Art. 33(2) and 33(3) PCT. Other claims are similarly recognized as such.

2) D3 (see respectively Figure 1 and text) disclose compounds falling into the wording of claim 55 as originally filed. An appropriate disclaimer of new claim 55 excludes now these compounds.

D2 discloses a compound which discloses some similarities with the compounds of claim 55 without being identical. Therefore claim 55 is new under Art. 33(2) PCT.

The problem underlying the present application is considered to reside in the provision of new preparations to modulate the immune response with AEP inhibitors and appropriate inhibitors which can be used to this purpose. The applicant has provided functional tests which demonstrate the effectiveness of such preparations, and new compounds as described in claim 55.

Therefore claim 1-56 are acceptable under Art. 33(3) PCT.

**INTERNATIONAL PRELIMINARY  
EXAMINATION REPORT - SEPARATE SHEET**

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International application No. PCT/GB99/00963

3) Claims 1-21 relate to subject-matter considered by this Authority to be covered by the provisions of Rule 67.1(iv) PCT. Consequently, no opinion will be formulated with respect to the industrial applicability of the subject-matter of these claims (Article 34(4)(a)(i) PCT).

**Re Item VIII**

**Certain observations on the international application**

As claim 55 finds support in the compounds of Table 1, it is found not to contravene the requirements of Art. 6 and Rule 6 PCT.

asparagine residue; and Q is a group capable of reacting with the active site of asparaginyl endopeptidase.

53. An inhibitor according to Claim 52 wherein the number of amino acid residues in (X<sub>a</sub>-X<sub>n</sub>) is between 1 and 25, preferably between 2 and 10.

54. An inhibitor according to Claim 53 which is any of BI-Ser-Gln-Asn-Q; BI-Leu-Glu-Asn-Q; BI-Leu-Gln-Asn-Q; BI-Pro-Glu-Asn-Q; BI-Leu-Lys-Asn-Q; BI-Gln-Asn-Q; BI-Glu-Asn-Q; BI-Asp-Glu-Asn-Q; BI-Asn-Gly-Asn-Q; BI-Phe-Pro-Asn-Q; BI-Val-Pro-Asn-Q; and BI-His-His-Asn-Q.

55. An inhibitor of asparaginyl endopeptidase which has the structure (X<sub>b</sub>-X<sub>c</sub>)Asn(X<sub>d</sub>-X<sub>e</sub>) wherein (X<sub>b</sub>-X<sub>c</sub>) are the r amino acid residues immediately N terminal to an Asn cleavage site in the invariant chain of Class II MHC molecules and (X<sub>d</sub>-X<sub>e</sub>) are the s amino acid residues immediately C terminal to an Asn cleavage site in the said invariant chain; Asn is an asparagine residue; and r and s are independently between 2 and 25, provided that the inhibitor is not the peptide CVFPNGTEVPNTRSRGHHN or the peptide ATKYGNMTEDHVMHLLQNA.

56. A composition comprising an inhibitor of asparaginyl endopeptidase and an inhibitor of cathepsin S.

EPO - DG 1

23. 03. 2000

(54)

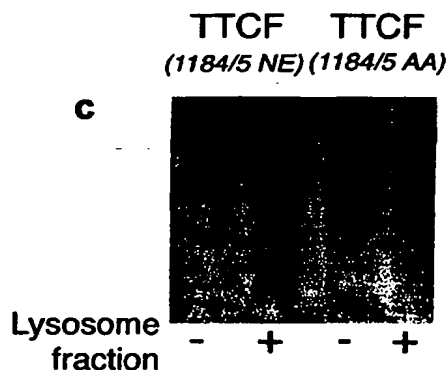
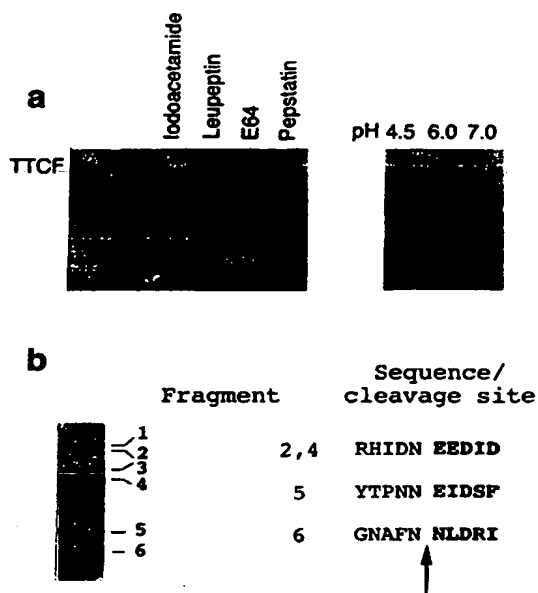
PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION  
International Bureau

## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

|   |  |   |   |
|---|--|---|---|
| (51) International Patent Classification <sup>6</sup> :<br><b>C07K 5/06, 5/08, A61K 38/55, C07K 14/705, G01N 33/68, A01K 67/027</b>   |  | <b>A1</b>   | (11) International Publication Number: <b>WO 99/48910</b>         |
|   |  |   | (43) International Publication Date: 30 September 1999 (30.09.99) |
| (21) International Application Number: - PCT/GB99/00963<br>(22) International Filing Date: 26 March 1999 (26.03.99)<br>(30) Priority Data:<br>9806442.1      26 March 1998 (26.03.98)      GB<br>60/086,966      28 May 1998 (28.05.98)      US<br>(71) Applicant (for all designated States except US): UNIVERSITY OF DUNDEE [GB/GB]; Dundee DD1 4HN (GB).<br>(72) Inventor; and<br>(75) Inventor/Applicant (for US only): WATTS, Colin [GB/GB]; 15 Westfield Terrace, Newport-on-Tay DD6 8HX (GB).<br>(74) Agent: MILES, John; Eric Potter Clarkson, Park View House, 58 The Ropewalk, Nottingham NG1 5DD (GB). |  | (81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).<br><br><b>Published</b><br><i>With international search report.</i><br><i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i> |   |

(54) Title: USE OF INHIBITORS OF MAMMALIAN ASPARAGINYL ENDOPEPTIDASE FOR THERAPY OF AUTOIMMUNE DISEASE



## (57) Abstract

A method of modulating the immune response in a patient in need of such modulation, the method comprising administering to the patient an effective amount of an inhibitor of asparaginyl endopeptidase. A method of reducing the processing of a protein antigen by a MHC Class II molecule by a cell, the method comprising contacting the cell with an inhibitor of asparaginyl endopeptidase.

**FOR THE PURPOSES OF INFORMATION ONLY**

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

|    |                          |    |                     |    |                       |    |                          |
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| CM | Cameroon                 |    | Republic of Korea   | PL | Poland                |    |                          |
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| CZ | Czech Republic           | LC | Saint Lucia         | RU | Russian Federation    |    |                          |
| DE | Germany                  | LI | Liechtenstein       | SD | Sudan                 |    |                          |
| DK | Denmark                  | LK | Sri Lanka           | SE | Sweden                |    |                          |
| EE | Estonia                  | LR | Liberia             | SG | Singapore             |    |                          |

USE OF INHIBITORS OF MAMMALIAN ASPARAGINYL ENDOPEPTIDASE FOR THERAPY  
OF AUTOIMMUNE DISEASE

The present invention relates to enzyme inhibition, and in particular to the inhibition of an enzyme that we call asparaginyl endopeptidase which we  
5 have found is involved in processing antigens, particularly microbial antigens, by the immune system.

The immune response to protein antigens involves a large number of individual gene products, and new ones are still being discovered. In  
10 some cases the gene products were known *per se*, but not known to be involved in the immune response.

An immune response may be raised towards foreign antigens, for example, antigens associated with microorganisms, or the immune  
15 response may be caused by a response to a self antigen (autoimmunity). In either case, there exists the need for methods and means for modulating the immune system and how it responds to these foreign and self antigens.

Because proteins must be proteolytically processed (ie partially degraded)  
20 before the T cells of the immune system can respond, one set of proteins of importance in an immune response are proteolytic enzymes (proteases). It is typically proteolysed proteins (ie protein fragments) which are presented by antigen presenting cells on MHC Class I or Class II molecules.

25

Several different aspartic and cysteine proteases are thought to be involved in invariant chain and antigen processing (see, for example, Fineschi & Miller (1997) *Trends Biochem. Sci.* 22, 377-382; and Chapman (1998)

*Curr. Op. Immunol.* 10, 93-102). For example, there is now good evidence that cathepsin S plays a role in the final stages of invariant chain processing (see, for example, Riese *et al* (1996) *Immunity* 4, 357-366; and Villadangos *et al* (1997) *J. Exp. Med.* 186, 549-560).

5

WO 97/40066 relates to the suppression of immune response by inhibiting cathepsin S.

We have now found, surprisingly, a further protease which is involved in processing protein antigens. The enzyme has asparaginyl endopeptidase (AEP) activity and, until the work described in this patent application, was not known to be involved in the immune response.

An enzyme with AEP activity was first detected in legumes and was called "legumain". Enzymes with AEP activity have also been detected in the blood fluke *Schistosoma mansoni* where it is involved in degrading haemoglobin (haemoglobinase) and, more recently, in mammals from which the cDNA has also been cloned (Chen *et al* (1997) *J. Biol. Chem.* 272, 8090-8096). For the avoidance of doubt, and in the event that different names are used, the animal enzymes with asparaginylendopeptidase activity may interchangeably be referred to as "asparaginyl endopeptidase", "AEP", "legumain" or "haemoglobinase". It will be appreciated that any particular animal, such as a human, may have more than one enzyme with asparaginyl endopeptidase activity, and that, in the context of the invention now described, an inhibitor of AEP may inhibit one or more forms of AEP.

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A first aspect of the invention provides a method of modulating the immune response in a patient in need of such modulation, the method comprising administering to the patient an effective amount of an inhibitor of asparaginyl endopeptidase.

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We have found that an enzyme with AEP activity is present in the lysosomes of B cells and is involved in degrading certain proteins. It is fragments of a protein, produced following proteolytic degradation, which are loaded onto and presented by Class II MHC molecules. Inhibition of  
10 AEP activity has been shown to interfere with the degradation of certain asparagine-containing proteins and to substantially inhibit the loading and presentation of peptides on Class II MHC molecule-containing cells (and to substantially inhibit T cell activation).

15 Furthermore, we have shown that AEP can degrade the so-called "invariant chain" which is a dedicated chaperone for class II MHC which must be removed by proteolysis before any peptide can be loaded (Cresswell (1996) *Cell* 84, 505). There are two alternatively spliced forms of the invariant chain, usually known as p31 and p41. Inhibition of  
20 AEP is believed to reduce the competency of Class II MHC molecules for binding antigenic peptides and reduce the presentation of antigenic peptides by Class II MHC molecules. Thus, inhibitors of AEP are likely to be immunosuppressive agents.

25 An enzyme with AEP activity is readily able to cleave the substrate Z-Ala-Ala-Asn-7-(4-methyl)coumarylamide to release a fluorescent product, where Z is benzyloxy carbonyl.

By an "inhibitor of asparaginyl endopeptidase" we include any suitable inhibitor. The inhibitor may be an enzyme inhibitor such as a competitive inhibitor or it may be a non-competitive inhibitor. The term also includes inhibitors of the provision of an active form of AEP in a cell such as  
5 inhibitors which inhibit the transcription of an AEP gene or the processing or translation of an AEP mRNA. As discussed below, suitable such inhibitors include AEP-selective antisense agents.

AEP is a cysteine protease and so inhibitors which chemically react with  
10 the active cysteine residue are suitable and are, typically, irreversible inhibitors. Other amino acid residues, such as histidine, may be present at the active site and inhibitors which react with any active site residue are suitable.

15 In one embodiment of the invention the AEP inhibitor is a competitive inhibitor. Typically, the competitive inhibitor is a peptide comprising an asparagine-containing peptide. Suitably, the peptide is a peptide comprising a known AEP cleavage site but which, because of its affinity for AEP, can compete for another AEP cleavage site and substantially  
20 prevent cleavage at the said other cleavage site if present in sufficient concentration. Known cleavage sites for AEP in tetanus toxin are described in Chen *et al* (1997) *J. Biol. Chem.* **272**, 8090-8098.

In a preferred embodiment of the invention the competitive inhibitor has  
25 the general structure  $(X_1)_p N(X_2)_q$  wherein  $X_1$  and  $X_2$  are amino acid residues, N is an asparagine residue, p is 3 to 6 and q is 1 to 3. Each of the p  $X_1$  residues is independently selected and each of the q  $X_2$  residues is independently selected.

Suitably, the competitive inhibitor is a peptide which comprises the peptide sequence Ala-Glu-Asn-Lys or, less preferably, Lys-Asn-Asn-Glu. Preferably, the competitive inhibitor is an N and C-terminal blocked peptide Ala-Glu-Asn-LysNH (AENK) or, less preferably, Lys-Asn-Asn-Glu-NH (KNNE).

As noted, and as is shown in Example 2, the class II MHC invariant chain is a substrate for AEP. Inspection of the invariant chain sequence indicates that any of the following asparagine residues may be processed by AEP: 71, 76, 131, 148 and 154 found in both p31 and p41 isoforms (p31 and p41 are two alternatively spliced forms of the invariant chain), and 217, 219, 238, 244 and 252 found only in the p41 isoform.

The sequence of the human invariant chain is given below. The p41 specific region is underlined.

MHRRRSRSCREDQKPVMDQDRLISNNEQLPMLGRRPGAPESKCSR GALYT  
GFSILVTLLLAGQATTAYFLYQQQGR LDKLTVTSQNLQLENLRMKLPKPPK  
PVSKMRMATPLLMQALPMGALPQGPMQNATKYGNMTEDHVMHLLQNADPLK  
VYPPLKGSFPENLTHLKNTMETIDWKVFESWMHWWLLFEMSRHSLEQKPTD  
APPKVLTKCQEEVSHIPAVHPGSFRPKCDENGNYLPLQCYGSIGYCWCVFP  
NGTEVPNTRSRGHHNCSESLELEDPS SGLGVTKQDLGPVPM

The complete sequence of the mRNA (and encoded polypeptide) for the HLA-DR-associated invariant chain is described in Strubin *et al* (1984) *EMBO J.* 3, 869-872. The structure of the human Ia-associated invariant (gamma)-chain is described in O'Sullivan *et al* (1986) *Proc. Natl. Acad. Sci. USA* 83, 4484-4488. Kudo *et al* (1985) *Nucl. Acids Res.* 13, 8827-

8841 describes the structure of the human gene encoding the invariant gamma chain of class II histocompatibility antigens.

Competitive inhibitors may be made based on the sequences which  
5 comprise, for example, two or three residues in each direction from the asparagine residue. Thus, for example, competitive inhibitors include residues 69 to 73, 68 to 74, 74 to 78, 73 to 79, 129 to 133, 128 to 134 and so on of the invariant chain. As is described in Example 2, both the p31 and p41 forms of the invariant chain are cleaved by AEP.

10 Figure 7 shows AEP processing of invariant chain.

Peptides may be synthesised by the Fmoc-polyamide mode of solid-phase peptide synthesis as disclosed by Lu *et al* (1981) *J. Org. Chem.* 46, 3433  
15 and references therein. Temporary N-amino group protection is afforded by the 9-fluorenylmethyloxycarbonyl (Fmoc) group. Repetitive cleavage of this highly base-labile protecting group is effected using 20% piperidine in N,N-dimethylformamide. Side-chain functionalities may be protected as their butyl ethers (in the case of serine threonine and tyrosine), butyl esters  
20 (in the case of glutamic acid and aspartic acid), butyloxycarbonyl derivative (in the case of lysine and histidine), trityl derivative (in the case of cysteine) and 4-methoxy-2,3,6-trimethylbenzenesulphonyl derivative (in the case of arginine). Where glutamine or asparagine are C-terminal residues, use is made of the 4,4'-dimethoxybenzhydryl group for protection of the side  
25 chain amido functionalities. The solid-phase support is based on a polydimethyl-acrylamide polymer constituted from the three monomers dimethylacrylamide (backbone-monomer), bisacryloylethylene diamine (cross linker) and acryloylsarcosine methyl ester (functionalising agent).

The peptide-to-resin cleavable linked agent used is the acid-labile 4-hydroxymethyl-phenoxyacetic acid derivative. All amino acid derivatives are added as their preformed symmetrical anhydride derivatives with the exception of asparagine and glutamine, which are added using a reversed N,N-dicyclohexyl-carbodiimide/1-hydroxybenzotriazole mediated coupling procedure. All coupling and deprotection reactions are monitored using ninhydrin, trinitrobenzene sulphonic acid or isotin test procedures. Upon completion of synthesis, peptides are cleaved from the resin support with concomitant removal of side-chain protecting groups by treatment with 95% trifluoroacetic acid containing a 50% scavenger mix. Scavengers commonly used are ethanedithiol, phenol, anisole and water, the exact choice depending on the constituent amino acids of the peptide being synthesised. Trifluoroacetic acid is removed by evaporation *in vacuo*, with subsequent trituration with diethyl ether affording the crude peptide. Any scavengers present are removed by a simple extraction procedure which on lyophilisation of the aqueous phase affords the crude peptide free of scavengers. Reagents for peptide synthesis are generally available from Calbiochem-Novabiochem (UK) Ltd, Nottingham NG7 2QJ, UK. Purification may be effected by any one, or a combination of, techniques such as size exclusion chromatography, ion-exchange chromatography and (principally) reverse-phase high performance liquid chromatography. Analysis of peptides may be carried out using thin layer chromatography, reverse-phase high performance liquid chromatography, amino-acid analysis after acid hydrolysis and by fast atom bombardment (FAB) mass spectrometric analysis.

In a further embodiment, the inhibitor may be a non-competitive or irreversible inhibitor. AEP activity can be blocked by high concentrations

of inhibitors which block all cysteine proteases and some serine proteases. However, it is particularly preferred that the inhibitors are selective for AEP. Non-competitive (irreversible or covalent) inhibitors are more preferred than competitive inhibitors.

5

Generally, irreversible AEP inhibitors may be any of peptide aldehydes, peptide halomethyl ketones (particularly peptide chloromethyl ketones and peptide fluoromethyl ketones), peptide diazomethanes (diazomethyl ketones) or peptide vinyl sulphones, peptide (acyloxy)methanes, peptide  
10 N, O diacyl hydroxamates, peptide nitriles, peptide  $\alpha$ -keto carbonyls, peptide ketomethylsulfonium salts, peptide epoxides or peptides which bind AEP and have a reactive group which will react with the active cysteine residue or other residue at the active site.

15 Suitably, the peptide sequence has the C-terminal residue as an asparagine residue to which is attached the group which reacts with the active site cysteine. The position of the asparagine residue in the peptide inhibitors (ie at the C terminus before the active group) is analogous to the P1 position at a cleavage site.

20

Thus, in general, an AEP inhibitor has the structure  $Bl-(X)_n-Asn-Q$  where Bl is a suitable N-terminal blocking group including, for example, acetyl or benzyloxycarbonyl; X is an amino acid residue; n is between 1 and 100, preferably between 1 and 50, more preferably between 1 and 10 and  
25 typically 2 to 6; Asn is an asparagine residue and Q is a group capable of reacting with the active site cysteine or other active site residue of AEP and, conveniently, forming a covalent complex thereby eliminating catalytic activity. It is preferred if the amino acid residues in the peptide

$X_n$ -Asn are in the L configuration and it is particularly preferred that, whether or not the  $X_n$  amino acid residues are in the L-configuration, that Asn is in the L-configuration. The amino acid residues X may be any naturally occurring amino acid residues or they may be residues which have non-natural side chains.

The peptide sequence before the C terminal asparagine and active group may be any suitable peptide sequence, but it will be appreciated that it is the sequence at a known AEP cleavage site. Cleavage sites for AEP are shown in, for example, Figures 7 and 8.

As noted, AEP cleaves both the p31 and p41 isoforms of the invariant chain. Non-competitive (or irreversible) inhibitors may readily be designed based on the residues N-terminal to the cleaved asparagine. As described above, any of the following asparagine residues may be processed by AEP: 71, 76, 131, 148 and 151 found in both p31 and p41, and 217, 219, 238, 244 and 252 found only in the p41 isoform. Thus, non-competitive inhibitors may be made which contain some of the amino acid residues N-terminal to these particular sites wherein the asparagine residue is linked to a group (Q) capable of reacting with the active site cysteine or other active site residue of AEP and, conveniently, forming a covalent complex thereby eliminating catalytic activity. Thus, AEP inhibitors may include BI-Ser-Gln-Asn-Q; BI-Leu-Glu-Asn-Q; BI-Leu-Gln-Asn-Q; BI-Pro-Glu-Asn-Q; BI-Leu-Lys-Asn-Q (corresponding to asparagines 71, 76, 131, 148 and 154).

Two preferred inhibitors are BI-Gln-Asn-Q and BI-Glu-Asn-Q.

Other AEP inhibitors may include BI-Asp-Glu-Asn-Q; BI-Asn-Gly-Asn-Q; BI-Phe-Pro-Asn-Q; BI-Val-Pro-Asn-Q; and BI-His-His-Asn-Q. These inhibitors relate to Asn residues of the p41 isoform with numbers 217, 219, 238, 244 and 252, respectively.

5

Other cleavage sites in unrelated proteins and peptides are shown in Figure 8. These may be used to design suitable inhibitors as described above.

10 Typically, peptide aldehyde inhibitors have the structure:

Acetyl (or other blocking group)-X<sub>n</sub>-Asparaginal

where X is an amino acid residue and n is between 1 and 100, preferably  
15 between 1 and 50, more preferably between 1 and 10 and typically 2 to 6.

A suitable peptide aldehyde includes acetyl-alanyl-glutamyl-asparaginal.

Elastatinal blocks AEP activity; however a more specific variant, in which  
20 the C terminal amino residue is replaced by asparagine, may be useful.

Figure 4 shows the structures of some useful inhibitors.

Typically, peptide chloromethylketone inhibitors have the structure.

25

Acetyl (or other blocking group)-X<sub>n</sub>-asparaginyl-chloromethyl ketone  
where X and n are as above.

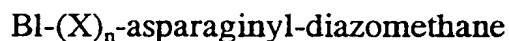


A suitable peptide includes acetyl-alanyl-glutamyl-asparaginyl-chloromethylketone and acetyl-tyrosyl-valyl-alanyl-asparaginyl-chloromethylketone (analogous to ICE protease inhibitor YVAD-cmk); see Figure 4 for further details.

5

Typically, peptide fluoromethylketone inhibitors have analogous structures to peptide chloromethylketone inhibitors except for the replacement of a chloro group with a fluoro group.

10 Typically, peptide diazomethane inhibitors have the structure

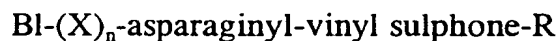


where Bl is any suitable blocking group including acetyl or benzyloxycarbonyl and X and n are as above. A diazomethane on the C-terminus of a peptide has a general structure  $\text{R-C(O)CHN}_2$  where R represents the peptide.

A suitable peptide includes Bl-alanyl-asparaginyl-diazomethane or Bl-alanyl-glutamyl-asparaginyl-diazomethane.

20

Typically, peptide vinyl sulphone inhibitors have the structure



25

where Bl is an N-terminal blocking group such as acetyl or benzyloxycarbonyl, X and n are as above, and R is any suitable alkyl or

aryl terminating group and includes C<sub>1</sub> to C<sub>10</sub> alkyl, phenyl, benzyl, naphthyl and the like.

A suitable peptide includes morpholinurea-leucyl-asparaginylyl-vinyl  
5 sulphone-phenyl or morpholinurea-alanyl-glutamyl-asparaginylyl-vinyl  
sulphone-phenyl.

Other suitable peptides include peptidyl(acyloxy) methanes and peptidyl  
N,O-diacylhydroxamates.

10

It will be appreciated from the foregoing that AEP inhibitors may be  
designed based on other cysteine protease inhibitors such as E-64 (1-trans-  
epoxysuccinyl-leucylamide (4-guanido)-butane), Leupeptin (acetyl-leucyl-  
leucyl-arginal), Antipain ([[(S)-1-carboxy-2-phenyl]-carbamoyl-Arg-Val-  
15 arginal), Elastinal (N-[(S)-1-carboxy-isopentyl]-carbamoyl- $\alpha$ -(2-  
iminohexahydro-4(S)-pyrimidyl]-L-glycyl-L-glutaminylyl-L-alaninal),  
TLCK (tosyllysylchloromethylketone) and TPCK  
(tosylphenylalanylchloromethylketone), for example by introduction of an  
asparaginylyl residue at an appropriate place in place of an existing amino  
20 acid residue.

Vinyl sulphone inhibitors of cysteine proteases are described in Palmer *et al* (1995) *J. Med. Chem.* **38**, 3193-3196. Peptide aldehyde inhibitors of  
cysteine proteases are described in Thomson *Methods Enzymol.* **46**, 220 -  
25 and Vinitsky *et al* (1994) *J. Biol. Chem.* **269**, 29860-29866 639-648.  
Peptide diazomethanes as inhibitors of cysteine proteases are described in  
Shaw (1994) *Methods Enzymol.* **244**, 649-656 and Shaw & Green (1981)  
*Methods Enzymol.* **80**, 820-826. Peptide (aryloxy) methanes as inhibitors

of cysteine proteases are described in Krantz (1994) *Methods Enzymol.* 244, 656-671. Peptide N,O-diarylhydroxamates as inhibitors of cysteine proteases are described in Brömme & Demuth (1994) *Methods Enzymol.* 244, 671-685. Peptide chloromethyl ketones as inhibitors of cysteine proteases are described in Williams & Mann (1993) *Methods Enzymol.* 222, 503-513. Thus, methods for synthesising suitable inhibitors are well known to the person skilled in the art.

It is preferred if the inhibitor is a chloromethylketone inhibitor or a fluoromethylketone inhibitor. It is less preferred if the inhibitor is a vinyl sulphone inhibitor. Chen *et al* (1998) *FEBS Letters* 441, 361-365 shows that legumain (AEP) has an active site similar to that of the proteases caspases, clostripain and gingipains. Caspases are aspartate-specific proteases whereas chlostripain and gingipains are arginyl and lysinyl proteases. AEP inhibitors may be made by modifying caspase inhibitors by replacing aspartate with asparagine, and they may be made by replacing arginine or lysine with asparagine in clostripain or gingipain inhibitors.

It is possible that, by analogy with other cysteine proteases, natural inhibitors of AEP exist, for example peptide inhibitors found in mammalian cells, or inhibitors found in microorganisms. The inhibitors for use in the invention include these inhibitors.

It is preferred that the AEP inhibitor is selective for AEP. By "selective" we mean that the inhibitor is at least ten times more potent for AEP than for another cysteine protease that does not have asparaginyl endopeptidase activity.

More preferably, the AEP inhibitor is at least 100 times more potent for AEP than for another cysteine protease inhibitor, still more preferably at least 1000 times more potent and most preferably at least 10 000 times more potent. In particular, it is preferred if the AEP inhibitor is an inhibitor which does not inhibit substantially cathepsins S and L. Cathepsins S and L are particular about residues which occupy the P2 position and inhibitors which contain aspartate or glutamate residues in the P2 position (ie the residue N-terminal of asparagine) are suitable for producing AEP inhibitors but are unfavourable for inhibitors of cathepsins L, S and B. Cathepsins S and L have a preference for uncharged amino acids in the P2 position (see Lysosomal Cysteine Proteases, Kirschke, Barrett & Rawlings, Second Edition, 1998, Oxford University Press, Oxford). Preferred inhibitors of AEP are those which include Ala-Glu-AsnQ where Q is defined as above. Particularly preferred inhibitors are those which include Ala-Glu-Asn-chloromethylketone or Ala-Glu-Asn-fluoromethylketone.

In a further preferred embodiment, the inhibitor of AEP may be an AEP-selective antisense agent.

Antisense agents are typically antisense oligonucleotides.

Antisense oligonucleotides are single-stranded nucleic acid, which can specifically bind to a complementary nucleic acid sequence. By binding to the appropriate target sequence, an RNA-RNA, a DNA-DNA, or RNA-DNA duplex is formed. These nucleic acids are often termed "antisense" because they are complementary to the sense or coding strand of the gene.

Recently, formation of a triple helix has proven possible where the oligonucleotide is bound to a DNA duplex. It was found that oligonucleotides could recognise sequences in the major groove of the DNA double helix. A triple helix was formed thereby. This suggests that it is possible to synthesise a sequence-specific molecules which specifically bind double-stranded DNA *via* recognition of major groove hydrogen binding sites.

By binding to the target nucleic acid, the above oligonucleotides can inhibit the function of the target nucleic acid. This could, for example, be a result of blocking the transcription, processing, poly(A)addition, replication, translation, or promoting inhibitory mechanisms of the cells, such as promoting RNA degradations.

Antisense oligonucleotides are prepared in the laboratory and then introduced into cells, for example by microinjection or uptake from the cell culture medium into the cells, or they are expressed in cells after transfection with plasmids or retroviruses or other vectors carrying an antisense gene. Antisense oligonucleotides were first discovered to inhibit viral replication or expression in cell culture for Rous sarcoma virus, vesicular stomatitis virus, herpes simplex virus type 1, simian virus and influenza virus. Since then, inhibition of mRNA translation by antisense oligonucleotides has been studied extensively in cell-free systems including rabbit reticulocyte lysates and wheat germ extracts. Inhibition of viral function by antisense oligonucleotides has been demonstrated *in vitro* using oligonucleotides which were complementary to the AIDS HIV retrovirus RNA (Goodchild, J. 1988 "Inhibition of Human Immunodeficiency Virus Replication by Antisense

Oligodeoxynucleotides", *Proc. Natl. Acad. Sci. (USA)* 85(15), 5507-11).

The Goodchild study showed that oligonucleotides that were most effective were complementary to the poly(A) signal; also effective were those targeted at the 5' end of the RNA, particularly the cap and 5' untranslated region, next to the primer binding site and at the primer binding site. The cap, 5' untranslated region, and poly(A) signal lie within the sequence repeated at the ends of retrovirus RNA (R region) and the oligonucleotides complementary to these may bind twice to the RNA.

Oligonucleotides are subject to being degraded or inactivated by cellular endogenous nucleases. To counter this problem, it is possible to use modified oligonucleotides, eg having altered internucleotide linkages, in which the naturally occurring phosphodiester linkages have been replaced with another linkage. For example, Agrawal *et al* (1988) *Proc. Natl. Acad. Sci. USA* 85, 7079-7083 showed increased inhibition in tissue culture of HIV-1 using oligonucleotide phosphoramidates and phosphorothioates. Sarin *et al* (1988) *Proc. Natl. Acad. Sci. USA* 85, 7448-7451 demonstrated increased inhibition of HIV-1 using oligonucleotide methylphosphonates. Agrawal *et al* (1989) *Proc. Natl. Acad. Sci. USA* 86, 7790-7794 showed inhibition of HIV-1 replication in both early-infected and chronically infected cell cultures, using nucleotide sequence-specific oligonucleotide phosphorothioates. Leither *et al* (1990) *Proc. Natl. Acad. Sci. USA* 87, 3430-3434 report inhibition in tissue culture of influenza virus replication by oligonucleotide phosphorothioates.

25

Oligonucleotides having artificial linkages have been shown to be resistant to degradation *in vivo*. For example, Shaw *et al* (1991) in *Nucleic Acids Res.* 19, 747-750, report that otherwise unmodified oligonucleotides become

more resistant to nucleases *in vivo* when they are blocked at the 3' end by certain capping structures and that uncapped oligonucleotide phosphorothioates are not degraded *in vivo*.

5 A detailed description of the H-phosphonate approach to synthesizing oligonucleoside phosphorothioates is provided in Agrawal and Tang (1990) *Tetrahedron Letters* 31, 7541-7544, the teachings of which are hereby incorporated herein by reference. Syntheses of oligonucleoside methylphosphonates, phosphorodithioates, phosphoramidates, phosphate  
10 esters, bridged phosphoramidates and bridge phosphorothioates are known in the art. See, for example, Agrawal and Goodchild (1987) *Tetrahedron Letters* 28, 3539; Nielsen *et al* (1988) *Tetrahedron Letters* 29, 2911; Jager *et al* (1988) *Biochemistry* 27, 7237; Uznanski *et al* (1987) *Tetrahedron Letters* 28, 3401; Bannwarth (1988) *Helv. Chim. Acta.* 71, 1517; Crosstick and Vyle (1989) *Tetrahedron Letters* 30, 4693; Agrawal *et al* (1990) *Proc.*  
15 *Natl. Acad. Sci. USA* 87, 1401-1405, the teachings of which are incorporated herein by reference. Other methods for synthesis or production also are possible. In a preferred embodiment the oligonucleotide is a deoxyribonucleic acid (DNA), although ribonucleic acid (RNA)  
20 sequences may also be synthesized and applied.

The oligonucleotides useful in the invention preferably are designed to resist degradation by endogenous nucleolytic enzymes. *In vivo* degradation of oligonucleotides produces oligonucleotide breakdown products of reduced  
25 length. Such breakdown products are more likely to engage in non-specific hybridization and are less likely to be effective, relative to their full-length counterparts. Thus, it is desirable to use oligonucleotides that are resistant to degradation in the body and which are able to reach the targeted cells.

The present oligonucleotides can be rendered more resistant to degradation *in vivo* by substituting one or more internal artificial internucleotide linkages for the native phosphodiester linkages, for example, by replacing phosphate with sulphur in the linkage. Examples of linkages that may be used include  
5 phosphorothioates, methylphosphonates, sulphone, sulphate, ketyl, phosphorodithioates, various phosphoramidates, phosphate esters, bridged phosphorothioates and bridged phosphoramidates. Such examples are illustrative, rather than limiting, since other internucleotide linkages are known in the art. See, for example, Cohen, (1990) *Trends in*  
10 *Biotechnology*. The synthesis of oligonucleotides having one or more of these linkages substituted for the phosphodiester internucleotide linkages is well known in the art, including synthetic pathways for producing oligonucleotides having mixed internucleotide linkages.

15 Oligonucleotides can be made resistant to extension by endogenous enzymes by "capping" or incorporating similar groups on the 5' or 3' terminal nucleotides. A reagent for capping is commercially available as Amino-Link II™ from Applied BioSystems Inc, Foster City, CA. Methods for capping are described, for example, by Shaw *et al* (1991) *Nucleic Acids*  
20 *Res.* **19**, 747-750 and Agrawal *et al* (1991) *Proc. Natl. Acad. Sci. USA* **88**(17), 7595-7599, the teachings of which are hereby incorporated herein by reference.

A further method of making oligonucleotides resistant to nuclease attack is  
25 for them to be "self-stabilized" as described by Tang *et al* (1993) *Nucl. Acids Res.* **21**, 2729-2735 incorporated herein by reference. Self-stabilized oligonucleotides have hairpin loop structures at their 3' ends, and show increased resistance to degradation by snake venom phosphodiesterase,



DNA polymerase I and fetal bovine serum. The self-stabilized region of the oligonucleotide does not interfere in hybridization with complementary nucleic acids, and pharmacokinetic and stability studies in mice have shown increased *in vivo* persistence of self-stabilized oligonucleotides with respect  
5 to their linear counterparts.

In accordance with the invention, the inherent binding specificity of antisense oligonucleotides characteristic of base pairing is enhanced by limiting the availability of the antisense compound to its intend locus *in*  
10 *vivo*, permitting lower dosages to be used and minimizing systemic effects. Thus, oligonucleotides are applied locally to achieve the desired effect. The concentration of the oligonucleotides at the desired locus is much higher than if the oligonucleotides were administered systemically, and the therapeutic effect can be achieved using a significantly lower total amount.  
15 The local high concentration of oligonucleotides enhances penetration of the targeted cells and effectively blocks translation of the target nucleic acid sequences.

The oligonucleotides can be delivered to the locus by any means appropriate  
20 for localized administration of a drug. For example, a solution of the oligonucleotides can be injected directly to the site or can be delivered by infusion using an infusion pump. The oligonucleotides also can be incorporated into an implantable device which when placed at the desired site, permits the oligonucleotides to be released into the surrounding locus.

25

The oligonucleotides may be administered *via* a hydrogel material. The hydrogel is noninflammatory and biodegradable. Many such materials now are known, including those made from natural and synthetic polymers. In a

preferred embodiment, the method exploits a hydrogel which is liquid below body temperature but gels to form a shape-retaining semisolid hydrogel at or near body temperature. Preferred hydrogel are polymers of ethylene oxide-propylene oxide repeating units. The properties of the polymer are dependent on the molecular weight of the polymer and the relative percentage of polyethylene oxide and polypropylene oxide in the polymer. Preferred hydrogels contain from about 10 to about 80% by weight ethylene oxide and from about 20 to about 90% by weight propylene oxide. A particularly preferred hydrogel contains about 70% polyethylene oxide and 30% polypropylene oxide. Hydrogels which can be used are available, for example, from BASF Corp., Parsippany, NJ, under the tradename Pluronic<sup>R</sup>.

In this embodiment, the hydrogel is cooled to a liquid state and the oligonucleotides are admixed into the liquid to a concentration of about 1 mg oligonucleotide per gram of hydrogel. The resulting mixture then is applied onto the surface to be treated, for example by spraying or painting during surgery or using a catheter or endoscopic procedures. As the polymer warms, it solidifies to form a gel, and the oligonucleotides diffuse out of the gel into the surrounding cells over a period of time defined by the exact composition of the gel.

The oligonucleotides can be administered by means of other implants that are commercially available or described in the scientific literature, including liposomes, microcapsules and implantable devices. For example, implants made of biodegradable materials such as polyanhydrides, polyorthoesters, polylactic acid and polyglycolic acid and copolymers thereof, collagen, and protein polymers, or non-biodegradable materials such as ethylenevinyl

acetate (EVAc), polyvinyl acetate, ethylene vinyl alcohol, and derivatives thereof can be used to locally deliver the oligonucleotides. The oligonucleotides can be incorporated into the material as it is polymerized or solidified, using melt or solvent evaporation techniques, or mechanically  
5 mixed with the material. In one embodiment, the oligonucleotides are mixed into or applied onto coatings for implantable devices such as dextran coated silica beads, stents, or catheters.

The dose of oligonucleotides is dependent on the size of the oligonucleotides  
10 and the purpose for which is it administered. In general, the range is calculated based on the surface area of tissue to be treated. The effective dose of oligonucleotide is somewhat dependent on the length and chemical composition of the oligonucleotide but is generally in the range of about 30 to 3000  $\mu\text{g}$  per square centimetre of tissue surface area.

15

The oligonucleotides may be administered to the patient systemically for both therapeutic and prophylactic purposes. The oligonucleotides may be administered by any effective method, for example, parenterally (eg intravenously, subcutaneously, intramuscularly) or by oral, nasal or other  
20 means which permit the oligonucleotides to access and circulate in the patient's bloodstream. Oligonucleotides administered systemically preferably are given in addition to locally administered oligonucleotides, but also have utility in the absence of local administration. A dosage in the range of from about 0.1 to about 10 grams per administration to an adult  
25 human generally will be effective for this purpose.

AEP-selective antisense agents, such as antisense oligonucleotides, may be designed and made by the person skilled in the art by reference, for

example, to the cDNA encoding AEP which is described in Chen *et al* (1997) *J. Biol. Chem.* 272, 8090-8098, incorporated herein by reference. It will be appreciated that there may be more than one AEP gene or that an AEP gene can give rise to more than one AEP transcript. Typically, the antisense oligonucleotide has at least 10 bases and, more preferably, at least 15 bases. Typically, the antisense oligonucleotides has around 18 to 20 bases. Although oligonucleotides may be designed and made which are complementary to any region of an AEP gene or mRNA transcript, in preferred embodiments the antisense oligonucleotides correspond to the region of the mRNA which contains the translation initiation signals or which encodes the N-terminus of AEP polypeptide. In addition, it is preferred if the sites in the AEP mRNA which are targeted by the antisense oligonucleotides are sites in which mRNA secondary structure is not expected and at which proteins are not expected to bind.

15

The AEP-inhibiting antisense agents need not be perfectly complementary to an AEP gene or mRNA in order to be effective. Rather, some degree of mismatches may be acceptable if the antisense oligonucleotide is of sufficient length. The oligonucleotides should have sufficient length and complementarity so as to selectively hybridise to an AEP transcript under physiological conditions. Preferably, mismatches are absent or minimal.

20

The AEP inhibitor, suitably, following administration to the patient, is able to contact and inhibit AEP at a suitable site in the patient. Although it is desirable for the AEP inhibitor to be cell permeant, it is believed that non-permeant AEP inhibitors may be useful since they may be taken into an appropriate cell by endocytosis.

25

An "effective amount" of the AEP inhibitor is an amount effective to modulate the immune response in the patient to a clinically useful extent.

AEP is believed to be involved in processing of protein antigens which are destined to be loaded into and presented by Class II MHC molecules. In a preferred embodiment of the invention the patient to be treated has or is at risk of a disease which involves MHC Class II molecules. Although we are not bound by any theory concerning the invention, we believe that AEP may play an important role in autoimmune disease because it may recognise and cleave at sites which are normally hidden by glycosylation at asparagine residues. Glycosylation at asparagine residues is common in mammalian proteins and glycosylated asparagine residues are not cleaved by AEP. An abnormal reduction in asparagine glycosylation of a self protein may lead to it being susceptible to AEP cleavage at these cryptic sites and, therefore, the peptide produced from the abnormally glycosylated protein may be susceptible to loading into and presentation to the immune system by Class II MHC molecules. Bacterial proteins, such as tetanus toxin, are typically not asparagine glycosylated and so, if they contain appropriate asparagine residues, are susceptible to degradation by AEP and presentation on Class II MHC molecules in any case.

Class II MHC molecules that remain associated with invariant chain cannot bind other peptides. Therefore, inhibition of AEP may reduce the competency of class II MHC molecules for binding antigenic peptides and reduces the presentation of antigenic peptides by Class II MHC molecules.

Preferably, the diseases are autoimmune diseases and it is particularly preferred if the patient to be treated has or is at risk of an autoimmune

disease such as rheumatoid arthritis, insulin-dependent diabetes mellitus, multiple sclerosis, Hashimoto's thyroiditis, coeliac disease, myasthenia gravis, pemphigus vulgaris, systemic lupus erythromatosus and Grave's disease.

5

In further preferred embodiments of the invention the patient to whom the inhibitor of AEP is administered has or is at risk of an allergic reaction or has undergone, or is to undergo, an organ or tissue or cell transplant in which graft rejection may ensue.

10

Several allergies and hypersensitivity reactions involve undesired responses to harmless environmental antigens inhaled, ingested or otherwise contacted. In several cases the antigens involved are known and, in as much as CD4 T cells are activated by class II MHC/peptide complexes, inhibition of processing of these antigens by AEP inhibitors is  
15 believed to be likely to block or reduce their display to the T cells which cause the allergic or hypersensitive reactions.

Typical allergies include:

20

Allergic rhinitis (hay fever) and allergic asthma induced by grass and tree pollens, dust-mite faeces, cat/dog fur and so on; contact hypersensitivity reactions induced by: metals (nickel, cobalt, copper, palladium), pentadecacatechol (poison ivy) or bee venom; food allergies to shellfish,  
25 milk, eggs, fish, wheat; and drug allergies eg to penicillin, aspirin.

Thus, the AEP inhibitors of the invention are useful in treating allergies and allergic reactions and hypersensitivity reactions. In addition to those

allergies and hypersensitivity reactions mentioned above, it is contemplated that the AEP inhibitors described herein are useful in treating allergic or hypersensitivity conditions associated with any of the following: the feline skin and salivary gland allergen of the domestic cat

5 *Felis domesticus* (the amino acid sequence of which is disclosed in WO 91/06571), the major protein allergens from the house dust mite dermatophagoides (amino acid sequences disclosed in WO 94/24281) and allergens present in any of the following: grass, tree and weed (including ragweed) pollens; fungi and molds; foods eg fish, shellfish, crab, lobster,

10 peanuts, nuts, egg and milk; stinging insects eg bee, wasp and hornet and the chironomidae (non-biting midges); spiders and mites; mammals such as dog, horse, rat, guinea pig, mice and gerbil; latex; biological detergent additives; drugs eg penicillins and other antibiotics and anaesthetic agents.

15 More particularly allergies to insect proteins arise from proteins from the housefly, fruit fly, sheep blow fly, screw worm fly, grain weevil, silkworm, honeybee, non-biting midge larvae, bee moth larvae, mealworm, cockroach and larvae of *Tenibrio molitor* beetle. All of these insect allergens are of particular relevance to allergic problems arising in

20 the workplace.

It will be appreciated that the AEP inhibitors may be used on their own, or that they may be administered in conjunction with other forms of treatment of the allergic or hypersensitivity conditions, which are known

25 in the art.

Allografts (from same species, different individual) or xenografts (different species) are frequently rejected because of differences at the

MHC genetic locus between the donor and the recipient. MHC matching improves the chances of graft acceptance as do immunosuppressive drug regimes. Blockade or reduction in the presentation of peptides on class II MHC molecules using the inhibitors of AEP described herein may be  
5 useful in immunosuppression and may be used instead of or in conjunction with existing suppressors of T cell activation such as cyclosporin or FK506.

Tissue and organ transplants which may benefit from additional  
10 immunosuppressive methods include kidney, lung, heart, liver, cornea and bone marrow. Prior treatment of both the organ and host may be beneficial.

It will be appreciated that the AEP inhibitors may be used on their own, or  
15 that they may be administered in conjunction with other forms of immunosuppression such as those described above or as mixtures of AEP inhibitors or as mixtures of an AEP inhibitor with, for example, an inhibitor of cathepsin S as described below.

20 The aforementioned inhibitors for use in the invention or a formulation thereof may be administered by any conventional method including oral, parenteral (eg subcutaneous or intramuscular) injection topical and the like. The treatment may consist of a single dose or a plurality of doses over a period of time.

25

Whilst it is possible for an inhibitor for use in the invention to be administered alone, it is preferable to present it as a pharmaceutical formulation, together with one or more acceptable carriers. The carrier(s)



must be "acceptable" in the sense of being compatible with the compound of the invention and not deleterious to the recipients thereof. Typically, the carriers will be water or saline which will be sterile and pyrogen free.

- 5 The formulations may conveniently be presented in unit dosage form and may be prepared by any of the methods well known in the art of pharmacy. Such methods include the step of bringing into association the active ingredient (inhibitor of AEP) with the carrier which constitutes one or more accessory ingredients. In general the formulations are prepared by
- 10 uniformly and intimately bringing into association the active ingredient with liquid carriers or finely divided solid carriers or both, and then, if necessary, shaping the product.

- Formulations in accordance with the present invention suitable for oral
- 15 administration may be presented as discrete units such as capsules, cachets or tablets, each containing a predetermined amount of the active ingredient; as a powder or granules; as a solution or a suspension in an aqueous liquid or a non-aqueous liquid; or as an oil-in-water liquid emulsion or a water-in-oil liquid emulsion. The active ingredient may also be presented as a bolus,
- 20 electuary or paste.

- A tablet may be made by compression or moulding, optionally with one or more accessory ingredients. Compressed tablets may be prepared by compressing in a suitable machine the active ingredient in a free-flowing
- 25 form such as a powder or granules, optionally mixed with a binder (eg povidone, gelatin, hydroxypropylmethyl cellulose), lubricant, inert diluent, preservative, disintegrant (eg sodium starch glycolate, cross-linked povidone, cross-linked sodium carboxymethyl cellulose), surface-active or

dispersing agent. Moulded tablets may be made by moulding in a suitable machine a mixture of the powdered compound moistened with an inert liquid diluent. The tablets may optionally be coated or scored and may be formulated so as to provide slow or controlled release of the active ingredient therein using, for example, hydroxypropylmethylcellulose in  
5 varying proportions to provide desired release profile.

Formulations suitable for topical administration in the mouth include lozenges comprising the active ingredient in a flavoured basis, usually  
10 sucrose and acacia or tragacanth; pastilles comprising the active ingredient in an inert basis such as gelatin and glycerin, or sucrose and acacia; and mouth-washes comprising the active ingredient in a suitable liquid carrier.

Formulations suitable for parenteral administration include aqueous and  
15 non-aqueous sterile injection solutions which may contain anti-oxidants, buffers, bacteriostats and solutes which render the formulation isotonic with the blood of the intended recipient; and aqueous and non-aqueous sterile suspensions which may include suspending agents and thickening agents. The formulations may be presented in unit-dose or multi-dose containers,  
20 for example sealed ampoules and vials, and may be stored in a freeze-dried (lyophilised) condition requiring only the addition of the sterile liquid carrier, for example water for injections, immediately prior to use. Extemporaneous injection solutions and suspensions may be prepared from sterile powders, granules and tablets of the kind previously described.

25

Preferred unit dosage formulations are those containing a daily dose or unit, daily sub-dose or an appropriate fraction thereof, of an active ingredient.

It should be understood that in addition to the ingredients particularly mentioned above the formulations of this invention may include other agents conventional in the art having regard to the type of formulation in question, for example those suitable for oral administration may include flavouring agents.

It will be appreciated that the pharmaceutical composition of the invention may further comprise an agent which is usefully administered to the patient in need of modulation of their immune response. For example, the composition may further comprise an agent for treatment or prevention or amelioration of an autoimmune disease or allergic or hypersensitivity reaction. As a further example, the composition may further comprise an immunosuppressive agent such as cyclosporin or FK506.

It is particularly preferred if the AEP inhibitor is administered at or near to the site of the disease. For example, in the treatment of rheumatoid arthritis it is desirable for the AEP inhibitor to be administered at or close to the joints affected. In this case, the AEP inhibitor may be administered by topical application.

A further aspect of the invention provides the use of an inhibitor of asparaginyl endopeptidase in the manufacture of a medicament for modulating the immune response in a patient in need of such modulation.

A still further aspect of the invention provides use of an inhibitor of asparaginyl endopeptidase for modulating the immune response in a patient in need of such modulation.

Yet still further aspects of the invention provide an inhibitor of asparaginyl endopeptidase for use in medicine and a pharmaceutical composition comprising an inhibitor of AEP and a pharmaceutically acceptable carrier.

- 5 In addition to AEP (as described for the first time herein), two other classes of proteases are involved in the Class II MHC pathway. These are the papain family of cysteine proteases (cathepsins S, L, B and H), and the aspartic acid family (cathepsins D and E).
- 10 Recent results from cathepsin L, D and B knock-out mice suggest that they have no major effect, at least in peripheral antigen presenting cells. This suggests that either there is considerable redundancy or that these enzymes are not the really important enzymes. The results described herein show that AEP is an important enzyme in processing Class II MHC
- 15 antigens, and it is contemplated that the inhibitors of AEP when used alone may be useful. However, in some circumstances it may be desirable to administer to the patient, or combine in a pharmaceutical formulation or in a therapeutic system (kit of parts), an inhibitor of AEP with an inhibitor of one or more of these other enzymes which are involved in the Class II
- 20 MHC pathway. In particular, since cathepsin S appears to be an important enzyme in invariant chain processing (see, for example, WO 97/40066) it is believed that a combination of cathepsin S inhibitors and AEP inhibitors may be particularly useful.
- 25 Thus, combinations of AEP inhibitors as disclosed herein and inhibitors cathepsin D or cathepsin E are contemplated. Known inhibitors of aspartic acid enzymes such as cathepsin D and cathepsin E include, for

example, pepstatin and CGP 53437 (Ciba-Geigy; see Alteri *et al* (1993) *Antimic. Ag. Chemotherapy* 37, 2087-).

Similarly, combinations of AEP inhibitors and lysosomal cysteine protease inhibitors (ie inhibitors of cathepsin S, L, B and H) are contemplated. Broad specificity inhibitors of these enzymes include leupeptin and the epoxide compound E64 or E64D. However, it is contemplated that cathepsin S inhibitors (in particular cathepsin S-selective or specific inhibitors) may be particularly advantageously combined with AEP inhibitors. Cathepsin S inhibitors are described in detail in WO 97/40066, all of which are incorporated herein by reference. In particular, the following cathepsin S inhibitors are believed to be useful in combination with AEP inhibitors: morpholino-Leu-homophenylalanine-vinyl sulphone; Leu-Leu-Leu-vinyl sulphone with various N-terminal blocking groups; vinyl sulphones based on the invariant chain cleavage sites cut by cathepsin S.

It is preferred that combinations of invariant chain-based inhibitors targeted to both AEP and cathepsin S are used. Combinations of such inhibitors may readily be devised by the skilled person by reference to the AEP inhibitors disclosed herein and by reference to the invariant chain-based cathepsin S inhibitors described in WO 97/40066. Examples of such combinations include, but are not limited to, BI-Glu-Asn-Leu-vinyl sulphone and/or BI-Leu-Arg-Met-vinyl sulphone (to block cathepsin S) plus one or more of BI-Ser-Gln-Asn; BI-Leu-Glu-Asn; BI-Leu-Gln-Asn; BI-Pro-Glu-Asn; and BI-Leu-Lys-Asn, all as vinyl sulphones (attached via the Asn residue).

As an alternative to vinyl sulphones as the reactive group, any of the reactive groups mentioned above ("Q") may be used in the inhibitors.

Other suitable combinations include invariant chain sequence based competitive inhibitors (as disclosed herein for AEP and as disclosed in  
5 WO 97/40066 for cathepsin S); and compounds disclosed herein as AEP inhibitors which are not based on the invariant chain in combination with any cathepsin S inhibitor.

10 The invention also includes a method of reducing the processing of a protein antigen by a MHC Class II molecule by a cell, the method comprising contacting the cell with an inhibitor of asparaginyl endopeptidase.

15 In one embodiment of the invention, an appropriate antigen presenting cell (APC), such as a dendritic cell or macrophage or B lymphocyte is transfected with a pool of cDNAs (or is contacted with proteins expressed from a pool of cDNAs) wherein the pool of cDNAs is believed to contain a cDNA encoding a protein which activates a T cell and may be associated  
20 with an autoimmune disease. The cDNA which encodes a protein which activates a T cell is selected and encodes a putative autoimmune protein. The assay may be carried out in the presence or absence of AEP inhibition in order to determine the involvement of AEP processing in the disease and also to determine any cryptic Asn glycosylation sites which may be  
25 important in the self-antigen immune response. Thus, the T cell is used to "read out" whether the cDNA for a putative autoantigen is being expressed and being presented, and whether AEP inhibitors have any effect on this.

A further aspect of the invention provides a method of identifying a compound for modulating Class II MHC antigen processing the method comprising contacting a test compound with asparaginyl endopeptidase and  
5 selecting a compound which reduces its activity.

This method (or screening assay) of the invention is suitable carried out in a format which allows for many test compounds to be screened simultaneously such as in a 96-well plate format. The activity of AEP is  
10 conveniently measured using any suitable substrate but it is preferred if the substrate is one which, upon cleavage by AEP, gives rise to a readily-detectable product. For example, the product may be coloured or fluorescent or detectable in some other way. It is particularly preferred if the product is fluorescent; a preferred substrate is Z-Ala-Ala-Asn-7-(4-  
15 methyl)coumarylamide.

The test compound is preferably a compound from a library of test compounds. The compound may be one which has been made by synthetic chemistry methods (such as combinatorial chemistry methods) or  
20 it may be a naturally occurring compound. Without prejudice to the type of compounds which can usefully be screened (which include any suitable compounds), it is believed that many inhibitors of AEP will have a peptide or peptide-like structure. Thus, it is preferred if the test compound has a peptide or peptide-like structure.

25

It will be appreciated that the method or screening assay may also include a step for eliminating non-specific inhibitory compounds or for positively

selecting those inhibitors which inhibit AEP but do not substantially inhibit other cysteine proteases which do not have AEP activity.

In a preferred embodiment of the invention, any test compounds which are identified as being capable of reducing the activity of AEP are selected for a second screening step which involves assessing their capacity to influence Class II MHC molecule loading and presentation in a suitable cell.

Thus, preferably the second step is the determination of whether the compound is capable of substantially inhibiting the loading and presentation of peptides on an appropriate Class II MHC molecule-containing cell. A convenient way of determining this is to assess whether the compound is capable of substantially inhibiting T cell activation by an appropriate Class II MHC molecule-containing cell. This may be done as described in the Example.

A further aspect of the invention comprises a non-human transgenic animal wherein a gene encoding asparaginyl endopeptidase has been modified and the animal expresses substantially no asparaginyl endopeptidase from said gene.

By "transgenic" animal we specifically include animals in which all or part of a gene have been "knocked out" or otherwise made substantially incapable of expressing an asparaginyl endopeptidase. Suitable mice can be made using standard methodology involving, for example, genetic manipulation of embryonic stem (ES) cells as is well known in the art. A cDNA encoding AEP is described in Chen *et al* (1997) *J. Biol. Chem.*



272, 8090-8096, and this information may be used in designing and making the transgenic animals of the invention.

It is particularly preferred if the non-human animal is a mouse, but it may  
5 be any suitable non-human animal including rat, rabbit and the like.

In a preferred embodiment of the invention, the non-human transgenic animal which expresses substantially no AEP from an AEP gene comprises a genetic background which predisposes the animal to an  
10 autoimmune disease either spontaneously or upon administration of protein antigen. Animals, especially mice, with a suitable genetic background are well known, for example NOD mice, which are a model for diabetes, EAE (experimental autoimmune encephalomyelitis) mice which are a model for allergic encephalomyelitis and CIA mice which are a model for  
15 collagen-induced arthritis. The mice used for the model disease EAE are of the MHC haplotypes H-2u or H-2k (eg PL/J or B10.PL strains). The mice used for the model disease CIA are of the MHC haplotype H-2q (eg DBA/1J strain). Such mice are described or referenced in Cantorna *et al* (1998) *J. Nutr.* 128, 68-72; Xiao & Link (1997) *Clin. Immunol.*  
20 *Immunopathol.* 85, 119-128; and Kumar *et al* (1997) *J. Exp. Med.* 185, 1725-1733.

Alternatively, the animal will be transgenic for a human Class II MHC molecule that predisposes to a human autoimmune disease.

25

In a further preferred embodiment, the non-human transgenic animal which expresses substantially no AEP from an AEP gene is further transgenic for a human Class II MHC molecule and, optionally, further

transgenic for CD4. Mice with these further characteristics are described in Altmann *et al* (1995) *J. Exp. Med.* 181, 867-875; Yamamoto *et al* (1994) *J. Ex. Med.* 180, 165-171; and Woods *et al* (1994) *J. Exp. Med.* 180, 173-181.

5

It will be appreciated that non-human animals which have combinations of characteristics can be made by crossing appropriate animals. Thus, the animals of the preferred embodiments of the invention may be made by crossing suitable animals.

10

From the foregoing it will be appreciated that new AEP inhibitors are disclosed.

15

A further aspect of the invention provides an inhibitor of asparaginyl endopeptidase which has the structure BI-(X<sub>a</sub>-X<sub>n</sub>)-Asn-Q wherein BI is any suitable N terminal blocking group; X<sub>a</sub>-X<sub>n</sub> are the n amino acid residues immediately N terminal to an Asn cleavage site in the invariant chain of Class II MHC molecules; Asn is an asparagine residue; and Q is a group capable of reacting with the active site of asparaginyl endopeptidase.

20

Suitable blocking groups, amino acids, and groups capable of reacting with the active site of AEP are the same as disclosed earlier in the application. It is preferred if the group capable of reacting with the active site of AEP can react with the active site cysteine residue.

25

The n amino acid residues N terminal to an Asn cleavage site can readily be determined by reference to the sequence of the invariant chain and by reference to the Asn residue number as described earlier. Typically the

number of amino acid residues is between 1 and 25; preferably, it is between 2 and 10; more preferably between 2 and 5.

Preferred inhibitors include Bl-Ser-Gln-Asn-Q; Bl-Leu-Glu-Asn-Q; Bl-Leu-Gln-Asn-Q; Bl-Pro-Glu-Asn-Q; Bl-Leu-Lys-Asn-Q; Bl-Gln-Asn-Q; Bl-Glu-Asn-Q; Bl-Asp-Glu-Asn-Q; Bl-Asn-Gly-Asn-Q; Bl-Phe-Pro-Asn-Q; Bl-Val-Pro-Asn-Q; and Bl-His-His-Asn-Q where Q is any reactive group as defined earlier and Bl is any N-terminal blocking group as defined earlier.

10

A further aspect of the invention provides an inhibitor of asparaginyl endopeptidase which has the structure (Xb-Xc)Asn(Xd-Xe) wherein (Xb-Xc) are the r amino acid residues immediately N terminal to an Asn cleavage site in the invariant chain of Class II MHC molecules and (Xd-Xe) are the s amino acid residues immediately C terminal to an Asn cleavage site in the said invariant chain; Asn is an asparagine residue; and r and s are independently between 2 and 25.

15

Preferably, r and s are independently between 2 and 10; more preferably r and s are independently between 2 and 5.

20

The invention will now be described by reference to the following Examples and Figures wherein

**Figure 1:** Processing of TTCF by a leupeptin-insensitive cysteine endopeptidase activity. (a) TTCF was digested *in vitro* in the presence of disrupted lysosomes (1-2  $\mu$ g) in the presence or absence of iodoacetamide (1 mM), leupeptin, E64 or pepstatin (each at 0.1 mg/ml). Digestions

25

were in 0.1M NH<sub>4</sub> acetate at pH 4.5 or Na<sub>2</sub>HPO<sub>4</sub>, pH 6.0 or 7.0. After 4 hours at 37°C, the reactions were separated by Tris-tricine SDS-PAGE and stained with Coomassie Blue. (b) Cleavage sites: TTCF digestion products were separated by SDS PAGE and electrophoretically transferred to nitrocellulose membrane. Individual fragments numbered 1-6 were subjected to 5 cycles of Edman degradation. N-terminal sequence obtained is shown in bold following upstream residues from TTCF (tetanus toxin numbering). The N-terminal sequence of His-tagged TTCF was obtained for fragments 1 and 3. (c) Residues Asn 1184 and Glu 1184 in TTCF were mutated to alanine (see Methods) and digestions performed as before. The mutation abolishes cleavage at the mutated site but not at the first site (doublet at 47 kD). In this experiment cleavage of the third site was minimal.

**Figure 2:** TTCF is processed by an asparaginyl endopeptidase. (a) Chromatography of the TTCF processing activity on Mono-S resin. Fractions eluted by NaCl gradient were incubated with the substrates Z-Ala-Ala-Asn-NHMec or Z-Phe-Arg-NHMec in the presence or absence of E64 (trans-epoxysuccinyl-L-leucylamido-(4-guanidino)butane). E64 had no inhibitory effect on the cleavage of Z-Ala-Ala-Asn-NHMec (not shown). (b) Fractions as above were incubated with 10 µg TTCF in the presence or absence of E64 for 4 hours at 37°C prior to SDS-PAGE analysis. (c) Each fraction was analysed by SDS-PAGE and Western blotting. Electrophoretically transferred material was probed with an affinity purified antiserum raised against the peptide KGIGSGKVLKSGPQC from human legumain<sup>1</sup>.

**Figure 3:** Peptide inhibitors of AEP block asparaginyl endopeptidase activity and TTCF processing *in vitro* and *in vivo*. (a) TTCF was digested with purified AEP (legumain from pig kidney) in the presence or absence of 0.2 mg/ml of the peptides indicated. Peptides were N and C-terminally blocked by not removing Fmoc or amide groups respectively. (b) Purified AEP or a crude lysosomal protease mixture were incubated with Z-Ala-Ala-Asn-NHMec, Z-Phe-Arg-NHMec (cathepsins L&B) or Z-Val-Val-Ala NHMec (cathepsin S) in the presence of increasing concentrations of AENK or AEQK. Release of 7-amino-4-methyl coumarin was measured after 10, 20 or 30 minutes. (c) Peripheral blood mononuclear cells were isolated from donor A.K. and preincubated for 15 minutes in the presence or absence of 1 mg/ml AENK (■) or AEQK (○) prior to addition of TTCF antigen. At different times the cells were washed, fixed in 0.05% glutaraldehyde and co-cultured with autologous TTCF specific T cell clones. (d) Human B-cell AEP digests TTCF and is inhibited by AENK but not AEQK.

**Figure 4** shows the structure of various AEP inhibitors.

**Figure 5** shows the structure of various protease inhibitors which can be converted to AEP inhibitors as described in the text.

**Figure 6** shows that AEP can cleave the Class II MHC invariant chain (both p31 and p41 isoforms).

**Figure 7** shows further details of AEP processing of the invariant chain.

Figure 8 shows the processing sites for AEP/legumain in diverse proteins. Proteins obtained from commercial sources, as described in the literature or custom peptides were digested with purified pig kidney legumain at pH 5.5 in the presence of 1 mM DTT. Digestion products were analysed by  
5 MALDI-TOF mass spectrometry and/or N-terminal Edman degradation following electrophoresis and electrophoretic transfer to PVDF membrane.

Figure 9 shows that TTCF processing and presentation requires AEP *in vivo*.  
10 a & b, Peptide inhibitors of asparaginyl endopeptidase slow antigen presentation *in vivo*. a, Peripheral blood mononuclear cells or b, EBV-transformed B cells, both from donor A.K. were preincubated for 15 minutes in the presence of AENK (■), AEQK (○) or no peptide (▲), prior to addition of TTCF antigen. When present peptide was maintained  
15 during antigen pulsing at 1 mg/ml (a) or 2 mg/ml (b). At different times the cells were washed, fixed in 0.05% glutaraldehyde and antigen presentation measured using proliferation of different autologous TTCF-specific T cell clones. c, AEP pre-processing of TTCF speeds antigen presentation. Antigen presentation was measured as in a, but using either  
20 TTCF (○) or TTCF pre-digested with AEP (▲, Δ) using either live (○, ▲) or fixed (Δ) autologous PBMC. d, Presentation of AEP-pre-processed antigen is no longer inhibited by AENK. Presentation was measured after 60 minutes of antigen pulsing using EBV B cells.

**Example 1: Asparaginyl endopeptidase is involved in processing and class II MHC presentation of a microbial antigen**

***Summary***

5

Foreign antigens must be proteolytically processed to allow loading of peptides onto class II MHC molecules. To investigate which proteases might be involved we exposed a domain of the microbial antigen, tetanus toxin (TTCF) to disrupted lysosomes purified from a human B cell line. Surprisingly, the dominant processing activity was not one of the known lysosomal cathepsins but rather an asparagine specific cysteine endopeptidase which we call AEP. This enzyme appears similar if not identical to a mammalian homologue of legumain<sup>1</sup>, an asparaginyl endopeptidase found originally in plants and parasites<sup>2,3</sup>. We designed competitive peptide inhibitors of AEP which specifically block its asparaginyl endopeptidase activity and inhibit processing of TTCF *in vitro* and its presentation *in vivo* to T cells. Since N-glycosylation renders asparagine resistant to cleavage by AEP we suggest that this enzyme may represent a further example of the ability of the innate immune system to focus its attention on microbial non-self.

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To analyse processing of a foreign antigen without making prior assumptions about which enzymes are involved we exposed a 47 kD domain of the tetanus toxin antigen (TTCF) to disrupted lysosomes isolated from the human B cell line EDR. As shown in Fig 1a, the antigen was fragmented to produce a discrete series of products at pH 4.5 but not at pH 7.0 and minimally at pH 6.0, confirming the lysosomal origin of the protease(s) involved (Fig 1a). Surprisingly, we could not

25

inhibit lysosomal digestion of TTCF with leupeptin or E64(trans-epoxysuccinyl-L-leucylamido-(4-guanidino)butane), broad spectrum inhibitors of lysosomal cysteine proteases or with pepstatin, an inhibitor of the aspartic acid cathepsins E and D (Fig 1a). LHSV, a more specific inhibitor of cathepsin S<sup>4</sup> was also without effect (not shown). Nor could we reproduce the lysosomal TTCF digestion pattern or indeed generate any of these fragments using purified cathepsins L, S, B, D or E, all of which have been implicated in the class II MHC antigen processing pathway (reviewed in refs 5 & 6). Further inhibitor studies revealed however that the activity was sensitive to iodoacetamide (Fig 1a), to N-ethyl maleimide and to high concentrations of the diazomethane, Z-Phe-Phe-CHN<sub>2</sub> (not shown) indicating that one or more cysteine protease(s) were nonetheless involved.

We sequenced each major TTCF digestion products to gain more information on this processing activity. All visible fragments arose from 3 cleavages in the TTCF protein and each occurred after an asparagine residue (Fig 1b). Mutation of asparagine 1184 and glutamic acid 1185 to alanine completely abolished cleavage at this site (Fig 1c) indicating that one (presumably the asparagine) or both of these residues is crucial for processing.

These results suggested that B cell lysosomes contain one or more novel cysteine endopeptidase(s) with possible specificity for asparagine. Although not previously described in antigen presenting cells, an enzyme with similar properties called legumain has been found in the seeds of leguminous plants<sup>2,3</sup>, in *Schistosoma mansoni*<sup>3</sup> and recently in some mammalian tissues such as kidney and placenta<sup>1</sup>. We next partially



purified the TTCF processing activity from a crude B cell lysosome fraction to establish whether or not it was an asparaginyl endopeptidase. As shown in Figure 2b, a peak of TTCF processing activity eluted from a cation-exchange resin at 0.4M NaCl. This activity co-eluted precisely  
5 with an activity capable of cleaving the substrate Z-Ala-Ala-Asn-7-(4-methyl)coumarylamide substrate (Fig 2a). Another protease activity, presumably cathepsin L and/or B, capable of cleaving the substrate Z-Phe-Arg-NHMec, partially overlapped the asparaginyl endopeptidase activity (Fig 2a). However this activity was completely inhibited by E64 (Fig 2a)  
10 while both the TTCF and Z-Ala-Ala-Asn-NHMec processing activity were unaffected (Fig 2b) confirming that a novel cysteine endopeptidase was involved. To compare this activity to the recently described mammalian form of legumain we generated anti-peptide antisera to several regions of the mammalian preprolegumain sequence<sup>1</sup>. Blotting of each fraction with  
15 an antisera to residues 125-140 (see methods) revealed the presence of a protein with the same mobility (35 kD) as purified pig kidney legumain which co-eluted with the TTCF and peptide substrate processing activity (Fig 2c). Finally, exactly the same processing sites were recognised by the crude lysosomal fraction, the partially purified TTCF processing  
20 activity and purified pig kidney legumain (Figs 1b & 2b, ref 1 and data not shown). Thus, mammalian legumain or a closely related enzyme, is present in human B lymphocytes and *in vitro*, is the major enzyme responsible for processing this antigen. We propose the name AEP (Asparaginyl endopeptidase) for this enzyme to distinguish it from the  
25 plant enzyme.

We decided to test the possibility that high concentrations of asparagine containing peptides might act as competitive inhibitors. As shown in Fig

3, the N and C-terminally blocked tetrapeptide Ala-Glu-Asn-Lys-NH (AENK) substantially inhibited the processing of TTCF by AEP (Fig 3a). Lys-Asn-Asn-Glu-NH (KNNE) also inhibited but was less effective. Importantly, the same concentrations of the glutamine analogues AEQK and KQQE did not inhibit TTCF processing demonstrating specific competitive inhibition of AEP (Fig 3a). To establish that inhibition of AEP was specific, we measured the hydrolysis of different synthetic peptide substrates in the presence and absence of these tetrapeptides. As shown in Fig 3b, AENK completely blocked cleavage of Z-Val-Ala-Asn-NHMec by B cell lysosomal fractions but had no effect on the hydrolysis of the cathepsin L/B substrate, Z-Phe-Arg-NHMec or on the Cathepsin S substrate Z-Val-Val-Arg-HNMec.

If AEP plays a role in TTCF processing in antigen presenting cells then sufficiently high levels of the competing AENK peptide might be expected to interfere with normal class II MHC loading of TTCF epitopes. Following a 15 minute preincubation with or without tetrapeptides, freshly isolated peripheral blood mononuclear cells (PBMC) were incubated with TTCF antigen in the continued presence or absence of the peptides. At different times, the cells were washed, fixed and then co-cultured with different autologous T cell clones to assess expression of different peptide/MHC complexes. Presentation to most clones was detectable after 60 minutes of antigen pulsing in the absence of competing peptide but in the presence of AENK, presentation to some clones was either undetectable (AK 20 and AK 90; Fig 3b) or profoundly inhibited (eg AK 6). At later times there was some recovery of presentation of these epitopes as expected for a competitive inhibitor. Several controls ruled out the possibility that the inhibitory effect of the AENK competitor is

simply due to non-specific toxicity. First, presentation to other clones recognising other regions of TTCF was not affected by the AENK competitor (eg AK 111, 71 and 33: Fig 3c). Secondly, there was a clear differential sensitivity to AENK versus AEQK (Fig 3c), strongly  
5 suggesting that specific inhibition of AEP was responsible for the slowed kinetics of presentation. Thirdly, the inhibitory effects of AENK could be overridden by using TTCF pre-digested *in vitro* by AEP. In other words, the requirement for AEP *in vivo*, and hence the inhibitory effect of AENK, was by-passed by AEP cleavage *in vitro* (data not shown). Taken  
10 together our results reveal AEP as a new and highly specific processing activity in the class II MHC pathway. Preliminary studies show that AEP is present in a variety of other antigen presenting cell types.

Although we uncovered this processing activity using an antigen  
15 proteolysis assay, it is possible that it also plays a role in invariant chain processing as well. Splenocytes from cathepsin D gene targeted mice continued to produce invariant chain processing products even when all known cysteine protease activity were also suppressed by leupeptin<sup>7</sup>. Villadangos *et al* suggested that other non-cysteine proteases (besides  
20 cathepsin D) must be initiating Ii (invariant chain) cleavage. Cathepsin E is one possible candidate but so too is AEP since, although it is a cysteine protease, it is completely resistant to leupeptin (Fig 1a). Intact p31 Ii is readily cleaved by AEP *in vitro* and we are currently assessing the possibility that AENK also interferes with Ii processing. However, the  
25 differential effect of the AENK inhibitor on T cell clones and the fact that pre-digestion of antigen with AEP abolished the inhibitory effect of the AENK inhibitor argues that its primary inhibitory effect in our studies was on TTCF processing.

Can any special significance be attached to the existence of an asparaginyl endopeptidase in antigen presenting cells? One possibility is that microbial proteins might in general be better substrates than host proteins since the latter could resist AEP attack by N-glycosylation of particularly sensitive asparagine residues. It is very striking that while TTCF is an extremely good substrate for AEP, cleavage occurs at only 3 of its 47 asparagine residues. This suggests that there may be additional determinants of AEP specificity to be identified and that strategic N-glycosylation might give substantial protection. Thus the innate immune systems ability to bind microbial antigen through recognition of non-self molecular 'patterns', might be harnessed to an 'innate' bias towards microbial antigen processing as well. In addition, our data raise the possibility that disturbances of N-glycosylation might result in altered processing of self proteins by AEP leading to presentation of 'cryptic' epitopes with possible pathological consequences.

### *Methods*

*TTCF processing activity.* Lysosomal fractions from the human B cell line EDR were prepared on 27% Percoll density gradients as previously described<sup>8</sup> using <sup>125</sup>I-labelled transferrin (internalised for 30 minutes at 37°C) and  $\beta$ -hexosaminidase to identify endosomal/plasma membrane and lysosomal fractions respectively. The peak of lysosomes was collected by centrifugation at 40,000 g for 1 hour, removed from the underlying Percoll pellet, collected again at 50,000 g for 1 hour and stored frozen. For separation of leupeptin-insensitive cysteine protease activity,  $7.5 \times 10^9$  cells were homogenised in a ball-bearing homogenizer and nuclei and

unbroken cells removed by centrifugation at 2,000 g for 10 minutes. A membrane pellet was collected at 40,000 g and solubilised in 50 mM citrate buffer, pH 5.5 containing 0.1% CHAPS (3-([cholamidopropyl]dimethylammonio]-1-propane-sulphonate). The extract  
5 was centrifuged at 2,000 g for 20 minutes and then applied to a Mono-S column (Pharmacia). Fractions were eluted with a gradient of NaCl and monitored for TTCF processing activity.

*Proteins and peptides.* A histidine tagged derivative of the C terminal  
10 domain of tetanus toxin was prepared<sup>9,10</sup> and purified<sup>10</sup> as described. This protein has residues 872-1315 of the complete toxin (1-1315) preceded by the sequence MGHGHHHHHHHHHSSGHIEGRHI. Mutagenesis of residues 1184/5 was performed using template EH106 obtained by cloning the His-TTCF (Reference 10) as an XbaI/BamHI fragment into pSL1180  
15 (Pharmacia). Site directed mutagenesis of residues 1184/85 was performed according to the method of Mikaelian and Sergent<sup>11</sup> using the mutagenic primer CGC TAC ACT CCG AAC GCG GCG ATC GAT TCT TTC GTT and flanking primers M13 rev (AGCGGATAACAATTTTCACACAGGA) and M13 seq  
20 (GTAAAACGACGGCCAGT). After sequencing to confirm mutagenesis, recloned back into pET16B for expression. Tetrapeptides were synthesised using Fmoc chemistry leaving the C-terminus amidated and the N-terminus retaining the Fmoc group. LHVS was a kind gift from Hidde Ploegh, MIT.

25

*Antigen presentation.* Peripheral blood mononuclear cells were prepared by Ficoll/Paque centrifugation and used fresh. T cell clones specific for TTCF were established from donor A.K. according to<sup>12</sup>. Epitope mapping

was performed using a set of 88 peptides, 17 residues in length spanning the TTCF sequence (Chiron Mimotopes). Clones AK 6, 90, and 71 recognise the peptides 1235-1245, 1145-1156 and 950-961 respectively. Clones 111 and 33 recognise epitopes within the regions 1225-1276 and 1104-1153. The epitope recognised by clone 20 has not been mapped. Peptides AENK or AEQK were dissolved in water, made isotonic with NaCl and diluted into RPMI growth medium. T cell proliferation assays were performed essentially as described<sup>12,13</sup>. Briefly, following antigen pulsing (30  $\mu$ g/ml TTCF) with or without tetrapeptides (1 mg/ml) PBMC were washed in PBS and fixed for 45 seconds in 0.05% glutaraldehyde. Glycine was added to a final concentration of 0.1M and the cells washed 5 times in RPMI 1640 medium containing 1% FCS before co-culture with T cell clones in round-bottom 96-well microtitre plates. After 48 hours the cultures were pulsed with 1  $\mu$ Ci of <sup>3</sup>H-thymidine and harvested 16 hours later.

Further experiments showed that AEP plays a role in TTCF processing *in vivo*. Freshly isolated peripheral blood mononuclear cells (PBMC) were pre-incubated briefly with AENK, or the control peptide AEQK, and then pulsed with TTCF antigen in the continued presence of one or other peptide. At different times, the cells were washed, fixed and then co-cultured with different autologous T cell clones to assess expression of different peptide/MHC complexes. Presentation was detectable between 30 and 60 minutes of antigen pulsing in the presence of the control peptide (Fig 9a) but in the presence of AENK, presentation to most clones (eg AK 5, 90 & 6) was undetectable at these early times. We also generated an EBV-B cell line from donor AK and tested the effect of the tetrapeptides

on B cell antigen presentation to the same T cell clones. As shown in Figure 9b, the AENK peptide blocked antigen presentation by EBV-B cells very effectively. The control peptide AEQK, had virtually no effect on the kinetics of antigen presentation (Fig 9b). Thus AENK specifically inhibited antigen presentation by two different APC populations. Antigen presentation to all clones recovered to different extents after longer times of antigen pulsing. We suggest that this recovery is expected given that AENK inhibits by acting as a competitive substrate and is itself hydrolysed by asparaginyl endopeptidases (data not shown). Under these conditions it is not possible to block TCTP processing completely. We also observed that some clones, for example clone 33 were less sensitive to blockade of AEP (Fig 4a & b). Clonal variations such as these might be explained either by a differential requirement for AEP processing, dependent on the location of the T cell epitope or on the antigen dose/response of individual clones or on a combination of these factors. Consistent with the first scenario, we have observed that clones specific for the same peptide region do show similar sensitivities to AEP blockade. For example, clones 5 & 90 and a third clone (AK 20) all recognise the 1145-1156 peptide and show very similar sensitivity to AENK (Fig 9 and data not shown). However, as expected, there was also variation in the antigen dose/response among the clones we tested. Those requiring higher levels of antigen for half-maximal triggering generally showed slower kinetics of presentation and greater sensitivity to AEP blockade. We suspect that the apparent differential requirement for AEP processing shown by different clones is due to a combination of epitope location, clone 'affinity' and possibly other factors. Nonetheless, it is clear that presentation to most tetanus toxin cell epitopes tested to date is affected by AEP blockade. This is consistent with the idea that initial processing by AEP may be the

'key' which 'unlocks' the native protein for further processing. As a further test of this hypothesis, we assayed the kinetics of presentation of TTCF that had been pre-digested *in vitro* with AEP, reasoning that this might accelerate antigen presentation. As shown in Figure 9c, this was indeed the case. Cells incubated with TTCF pre-processed with AEP presented T cell epitopes after only 15-30 minutes whereas unprocessed TTCF required 60-120 minutes before presentation could be detected. Importantly, AENK did not block the rapid presentation of the pre-processed antigen at these times (Fig 9d), confirming that this inhibitor specifically blocks AEP action on the antigen substrate *in vivo*. TTCF pre-processed with AEP still required further cellular processing since it could not be presented by fixed PBMC (Fig 9c & d). Taken together these results support a working model whereby AEP initiates TTCF proteolysis but that generation of suitable peptides for MHC binding requires further intracellular processing. Native TTCF was a poor substrate for cathepsins E,D,L,S and B (data not shown) but many processing sites, for example for cathepsins E and D were revealed following antigen unfolding.

20

We also analysed a natural 21-residue tryptic glycopeptide (residues 622-642) purified from human serum transferrin (see Table 1). This peptide contains 2 asparagine residues one of which (N630) carries a biantennary oligosaccharide (Lu, J. & van Halbeek, H. (1996) "Complete <sup>1</sup>H and <sup>13</sup>C resonance assignments of a 21-amino acid glycopeptide prepared from human serum transferrin." *Carbohydr. Res.* **296**, 1-21). The same peptide was synthesised without modification of the asparagines and digested in parallel. [3 lysine residues were added to improve solubility].

25



Both mass spectrometric analysis and N-terminal sequencing of the products confirmed that hydrolysis of both asparagines 630 and 637 occurred in the non-glycosylated peptide (Table 1). However, we only detected products from hydrolysis of N637 in the glycopeptide (Table 1).

- 5 A second transferrin-derived tryptic glycopeptide (residues 421-452) containing 3 asparagine residues was also digested with AEP. Again, we only detected fragments from hydrolysis at the non-glycosylated asparagines 430 & 436 (not shown).

10 *Table 1: N-glycosylation blocks asparaginyl endopeptidase action on a natural glycopeptide*

|  | Predicted Mass                              | Observed Mass   |
|--|---|---|
| Synthetic Transferrin Peptide (622-642KKK)   |   |   |
| No enzyme<br>QQQHLFGSNVTDCSGNFCLFRKKK  | 2784.36                                     | 2784.17   |
| + Asparagine endopeptidase<br>QQQHLFGSN<br>VTDCSGNFCLFRKKK<br>FCLFRKKK                             | 1058.51<br>1745.83<br>1069.63               | 1058.46<br>1743.69*<br>1069.59                        |
| Transferrin glycopeptide (622-642)   |   |   |
| No Enzyme<br>QQQHLFGSN(CHO)VTDCSGNFCLFR  | 4720.76                                     | 4720.2  |
| + Asparagine endopeptidase<br><br>QQQHLFGSN(CHO)VTDCSGN<br>FCLFR<br>VTDCSGNFCLFR<br>QQQHLFGSN(CHO) | <br>3996.88<br>742.91<br>1476.66<br>3263.13 | <br>3996.85<br>742.23<br>not observed<br>not observed |

- 15 Predicted fragment masses were obtained from the human transferrin sequence (accession number: P02787) calculated using PeptideMass at the EXPASY server. The transferrin biantennary oligosaccharide (2 Sialic acid + 4 GlcNAc + 5 hexose has a mass of 2204 coupled to peptide Lu, J. & van Halbeek, H. (1996) "Complete <sup>1</sup>H and <sup>13</sup>C resonance

assignments of a 21-amino acid glycopeptide prepared from human serum transferrin." *Carbohydr. Res.* **296**, 1-21). Variants also observed but not shown are monosialylated forms of the glycopeptides and N-terminal peptide forms 17 mass units lower corresponding to N-terminal glutamine pyroglutamisation. \*Cyclisation of the two cysteine residues in this peptide likely accounts for the observed mass deficit of 2.

**Example 2: Asparaginyl endopeptidase can process the Class II MHC invariant chain**

10

A key step in the generation of peptide/class II MHC complexes is the complete removal of a protein called the invariant chain from the class II MHC  $\alpha\beta$  dimer. This is a dedicated chaperone for class II MHC which must be removed by proteolysis before any peptide can be loaded (Cresswell (1996) *Cell* **84**, 505-). Normally, its removal requires, first, proteolysis to produce a small fragment (known as CLIP ~about 3 kD) which still blocks the class II peptide binding site, and second, the action of a protein called DM which removes this peptide. Blocking invariant chain removal effectively shuts down loading of all other peptides. There are two alternatively spliced forms of the invariant chain usually known as p31 and p41. p41 is minor in most cells except dendritic cells where it accounts for around 40% of invariant chain.

20

*In vitro*, cathepsin S can produce the small fragment (CLIP) suitable for DM action (Riese *et al* (1996) *Immunity* **4**, 357-) but it is likely that other enzymes are involved in invariant chain processing in living cells. For example, inhibitors of cathepsin S result in the accumulation of an invariant chain fragment considerably larger (~ 10 kD) than CLIP. This

25

shows that cathepsin S is only really essential at the later stages of invariant chain processing and that other enzymes can produce the 10 kD fragment. It has now been found that AEP can degrade the invariant chain *in vitro*. Thus, AEP processing of the invariant chain may initiate its destruction in some or all cells (some cells, eg thymic cortical epithelial cells do not seem to express cathepsin S).

The human EBV-transformed lines PALA or AL-EBV were radiolabelled for 15 minutes at 37°C with 0.5 mCi/ml <sup>35</sup>S-Translabel (ICN), washed and lysed in Tris-buffered saline containing 1% Triton X-100 plus protease inhibitors (Davidson & Watts (1989) *J. Cell. Biol.* **109**, 85). The lysates were cleared by centrifugation and pre-cleared with Protein G sepharose. Invariant chain/class II MHC complexes were precipitated with the VIC-Y1 antibody (gift from W Knapp). The complexes were collected on Protein G sepharose, washed 3 times in Tris-buffered saline, 1% Triton X-100 and resuspended in 50 mM citrate buffer, 1 mM EDTA, 5 mM dithiothreitol with or without 0.05% CHAPS (Calbiochem). Purified aspariginyl endopeptidase was added to 5 mU per 20 µl reaction. Digestions were performed at 37°C for 30 minutes and analysed by Tris-tricine SDS-PAGE (12%). Note that both p41 and p31 forms of the invariant chain are completely digested to produce 3-4 discrete fragments (Figure 6) while both the α and β chains of class II MHC are resistant to AEP processing. The presence of the detergent CHAPS does not affect the digestion.

Circumstantial evidence has previously shown that enzymes, other than cathepsin S, L, B and D, can degrade the invariant chain in living cells. This comes from the finding that in spleen cells from a cathepsin D knock-

out mouse also treated with leupeptin (which inhibits cathepsins S, L and B), the early stages of invariant chain processing still proceed (Villadangos *et al* (1997) *J. Exp. Med.* 186, 549-). The results presented here suggest that AEP could be responsible for this residual invariant chain processing activity.

A further experiment was carried out to demonstrate invariant chain processing. The results are shown in Figure 7.

10 The human EBV-transformed cell line Pala was labelled with <sup>35</sup>S-Methionine/Cysteine (Translabel, ICN) for 30 minutes and the cells lysed for immunoprecipitation. Cleared lysates were incubated with either mAb DA6.231 (class II MHC  $\beta$ -chain specific) or VIC- $\gamma$ 1 (invariant chain specific). Recovered immunoprecipitates were then digested with 5 mU  
15 purified pig kidney legumain for 60 minutes. The digests were then run on 12% Tris-Tricine SDS gels, transferred to PVDF membrane and the numbered fragments in panel (a) of Figure 7 arising from AEP processing subjected to N-terminal Edman degradation. The eluate from each cycle of cleavage was dried down, resuspended in scintillation cocktail and  
20 counted on a  $\beta$  counter. Similar digestion patterns were obtained for the DA6.231 precipitates (not shown). <sup>35</sup>S cpm released at specific cycles of Edman degradation allowed the Asparagine residues cleaved to be identified. An illustrative example of the raw data obtained from sequencing band 5 is shown in (b) of Figure 7. Figure 7(c) Human li p31  
25 sequence showing 3 AEP processing sites. AEP can cleave on both sides of the CLIP region which is underlined. Note that the first site obtained from sequencing band 1 is in the cytoplasmic tail and would not be available for processing in living cells.

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**CLAIMS**

1. A method of modulating the immune response in a patient in need of such modulation, the method comprising administering to the patient an effective amount of an inhibitor of asparaginyl endopeptidase.  
5
2. A method according to Claim 1 wherein the patient has or is at risk of a disease which involves MHC Class II molecules.
- 10 3. A method according to Claim 1 or 2 wherein the disease is an autoimmune disease.
4. A method according to Claim 3 wherein the disease is rheumatoid arthritis.  
15
5. A method according to Claim 1 or 2 wherein the patient has or is at risk of an allergic or hypersensitivity reaction.
6. A method according to Claim 1 or 2 wherein the patient has undergone or is to undergo a transplant.  
20
7. A method according to Claim 6 wherein the material transplanted, or to be transplanted, has been contacted with an effective amount of an inhibitor of asparaginylendopeptidase.  
25
8. A method according to any one of the preceding claims wherein the inhibitor is a competitive inhibitor.

9. A method according to Claim 8 wherein the competitive inhibitor is a peptide comprising is an asparagine-containing peptide.

10. A method according to Claim 9 wherein the peptide is an N and C-terminal blocked peptide Ala-Glu-Asn-Lys-NH (AENK) or Lys-Asn-Asn-Glu-NH (KNNE).

11. A method according to Claim 1 to 6 wherein the inhibitor is a non-competitive or irreversible inhibitor.

12. A method according to Claim 11 wherein the inhibitor has the structure  $Bl-(X)_n-Asn-Q$  where Bl is any suitable N terminal blocking group; X is an amino acid residue; n is between 1 and 100, Asn is an asparagine residue and Q is a group capable of reacting with the active site cysteine of asparaginyl endopeptidase.

13. A method according to any one of the preceding claims further comprising administering to the patient an effective amount of an agent for treatment or prevention or amelioration of an autoimmune disease or an allergic or hypersensitivity reaction.

14. A method according to any one of Claims 1 to 12 further comprising administering to the patient an effective amount of an immunosuppressive agent.

15. A method of reducing the processing of a protein antigen by a MHC Class II molecule by a cell, the method comprising contacting the cell with an inhibitor of asparaginyl endopeptidase.



16. A method according to Claim 15 wherein the inhibitor is a competitive inhibitor.
- 5 17. A method according to Claim 16 wherein the competitive inhibitor is a peptide comprising an asparagine-containing peptide.
18. A method according to Claim 17 wherein the peptide is an N and C-terminal blocked peptide Ala-Glu-Asn-Lys-NH (AENK) or Lys-Asn-  
10 Asn-Glu-NH (KNNE).
19. A method according to Claim 15 wherein the inhibitor is a non-competitive or irreversible inhibitor.
- 15 20. A method according to Claim 19 wherein the inhibitor has the structure  $Bl-(X)_n-Asn-Q$  where Bl is any suitable N terminal blocking group; X is an amino acid residue; n is between 1 and 100, Asn is an asparagine residue and Q is a group capable of reacting with the active site cysteine of asparaginyl endopeptidase.
- 20 21. A method according to any one of Claims 15 to 20 wherein the cell is, or is comprised in a tissue or organ, for transplantation into a patient.
22. Use of an inhibitor of asparaginyl endopeptidase in the manufacture  
25 of a medicament for modulating the immune response in a patient in need of such modulation.

23. Use according to Claim 22 wherein the patient has or is at risk of a disease which involves MHC Class II molecules.

24. Use according to Claim 22 or 23 wherein the disease is an  
5 autoimmune disease.

25. Use according to Claim 24 wherein the disease is rheumatoid arthritis.

10 26. Use according to Claim 22 or 23 wherein the patient has or is at risk of an allergic or hypersensitivity reaction.

27. Use according to Claim 22 or 23 wherein the patient has undergone or is to undergo a transplant.

15

28. Use according to any one of Claims 22 to 27 wherein the inhibitor is a competitive inhibitor.

29. Use according to Claim 28 wherein the competitive inhibitor is a  
20 peptide comprising is an asparagine-containing peptide.

30. Use according to Claim 29 wherein the peptide is an N and C-terminal blocked peptide Ala-Glu-Asn-Lys-NH (AENK) or Lys-Asn-Asn-Glu-NH (KNNE).

25

31. Use according to any one of Claims 22 to 27 wherein the inhibitor is a non-competitive or irreversible inhibitor.

32. Use according to Claim 31 wherein the inhibitor has the structure  $Bl-(X)_n-Asn\ Q$  where Bl is any suitable N terminal blocking group; X is an amino acid residue; n is between 1 and 100, Asn is an asparagine residue and Q is a group capable of reacting with the active site cysteine of asparaginyl endopeptidase.

33. Use according to any one of Claims 22 to 32 wherein the patient is administered an effective amount of an agent for treatment or prevention or amelioration of an autoimmune disease or an allergic or hypersensitivity reaction.

34. Use according to any one of Claims 22 to 32 wherein the patient is administered an effective amount of an immunosuppressive agent.

35. Use of an inhibitor of asparaginyl endopeptidase for modulating the immune response in a patient in need of such modulation.

36. Use of an inhibitor of asparaginyl endopeptidase for reducing the processing of a protein antigen by a MHC Class II molecule by a cell.

37. An inhibitor of asparaginyl endopeptidase for use in medicine.

38. A pharmaceutical composition comprising an inhibitor of asparaginyl endopeptidase and a pharmaceutically acceptable carrier.

39. A pharmaceutical composition according to Claim 38 further comprising an agent which is usefully administered to a patient in need of modulation of the immune response.

40. A pharmaceutical composition according to Claim 38 further comprising an agent for treatment or prevention or amelioration of an autoimmune disease or an allergic or hypersensitivity reaction.

5

41. A pharmaceutical composition according to Claim 38 further comprising an immunosuppressive agent.

42. A pharmaceutical composition comprising an inhibitor of asparaginyl endopeptidase, an inhibitor of cathepsin S and a pharmaceutically acceptable carrier.

10

43. A method of identifying a compound for modulating Class II MHC antigen processing the method comprising contacting a test compound with asparaginyl endopeptidase and selecting a compound which reduces its activity.

15

44. A method according to Claim 43 wherein the activity of asparaginyl endopeptidase is measured using a substrate which upon cleavage by said endopeptidase, yields a readily detectable product.

20

45. A method according to Claim 44 wherein the substrate is Z-Ala-Ala-Asn-7-(4-methyl)coumarylamide and the product is fluorescent.

46. A method according to any one of Claims 43 to 45 the method further comprising the step of determining whether the so selected compound is capable of substantially inhibiting the loading and

25

presentation of peptides on an appropriate Class II MHC molecule-containing cell.

47. A method according to Claim 46 wherein it is determined whether  
5 the so selected compound is capable of substantially inhibiting T cell activation by an appropriate Class II MHC molecule-containing cell.

48. A non-human transgenic animal wherein a gene encoding asparaginyl endopeptidase has been modified and the animal expresses  
10 substantially no asparaginyl endopeptidase from said gene.

49. A non-human transgenic animal according to Claim 48 which is a mouse.

15 50. A non-human transgenic animal according to Claim 48 or 49 further comprising a genetic background which predisposes to an autoimmune disease either spontaneously or upon administration of protein antigen.

20 51. A non-human transgenic animal according to Claims 48 or 49 further transgenic for a human Class II MHC molecule and, optionally, further transgenic for human CD4.

52. An inhibitor of asparaginyl endopeptidase which has the structure  
25  $\text{Bl}-(\text{X}_a-\text{X}_n)-\text{Asn}-\text{Q}$  wherein Bl is any suitable N terminal blocking group;  $\text{X}_a-\text{X}_n$  are the n amino acid residues immediately N terminal to an Asn cleavage site in the invariant chain of Class II MHC molecules; Asn is an

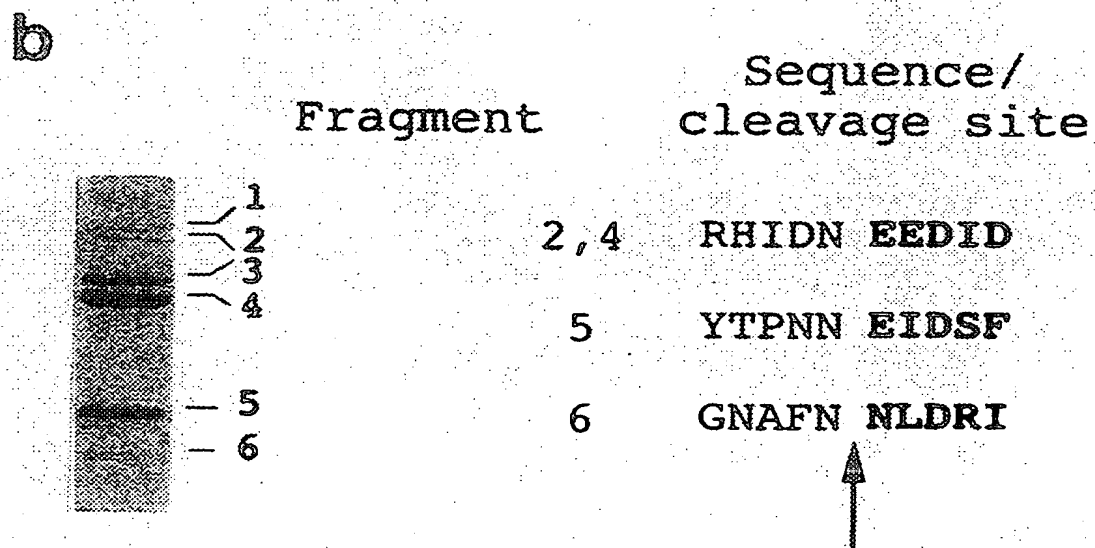
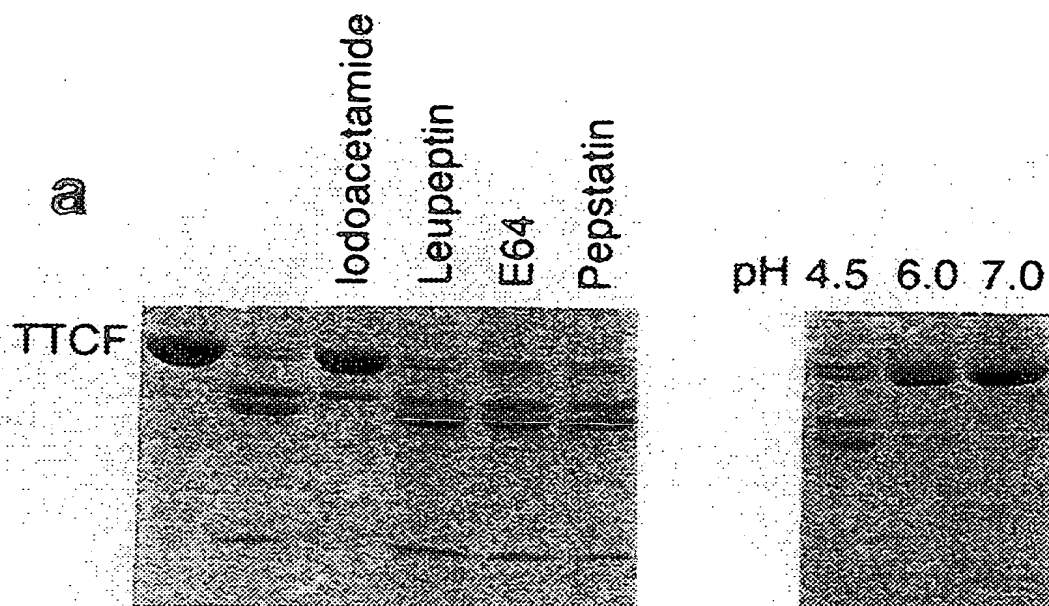
asparagine residue; and Q is a group capable of reacting with the active site of asparaginyl endopeptidase.

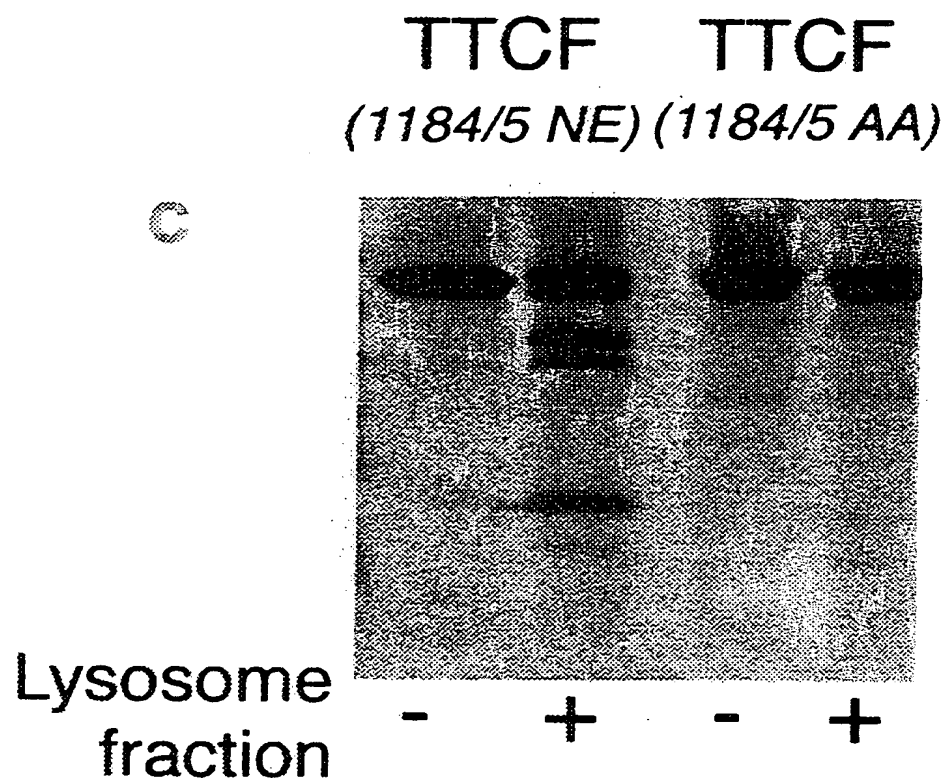
53. An inhibitor according to Claim 52 wherein the number of amino acid residues in  $(X_a-X_n)$  is between 1 and 25, preferably between 2 and 10.

54. An inhibitor according to Claim 53 which is any of Bl-Ser-Gln-Asn-Q; Bl-Leu-Glu-Asn-Q; Bl-Leu-Gln-Asn-Q; Bl-Pro-Glu-Asn-Q; Bl-Leu-Lys-Asn-Q; Bl-Gln-Asn-Q; Bl-Glu-Asn-Q; Bl-Asp-Glu-Asn-Q; Bl-Asn-Gly-Asn-Q; Bl-Phe-Pro-Asn-Q; Bl-Val-Pro-Asn-Q; and Bl-His-His-Asn-Q.

55. An inhibitor of asparaginyl endopeptidase which has the structure  $(X_b-X_c)\text{Asn}(X_d-X_e)$  wherein  $(X_b-X_c)$  are the  $r$  amino acid residues immediately N terminal to an Asn cleavage site in the invariant chain of Class II MHC molecules and  $(X_d-X_e)$  are the  $s$  amino acid residues immediately C terminal to an Asn cleavage site in the said invariant chain; Asn is an asparagine residue; and  $r$  and  $s$  are independently between 2 and 25.

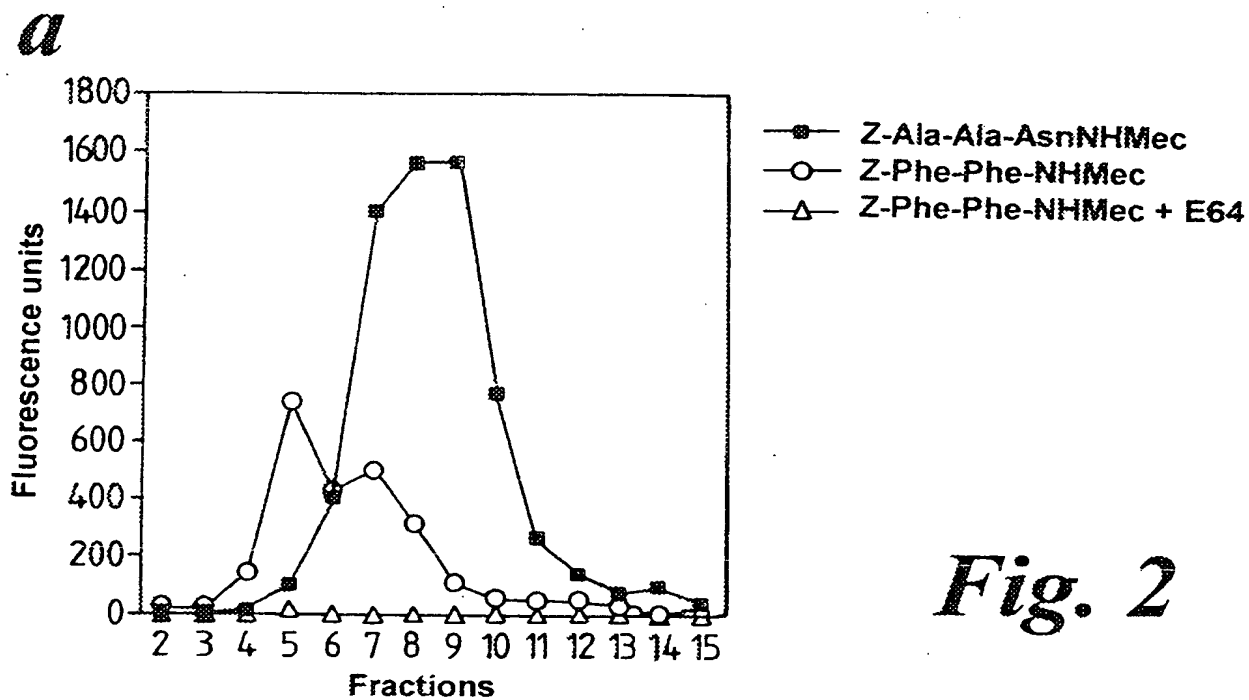
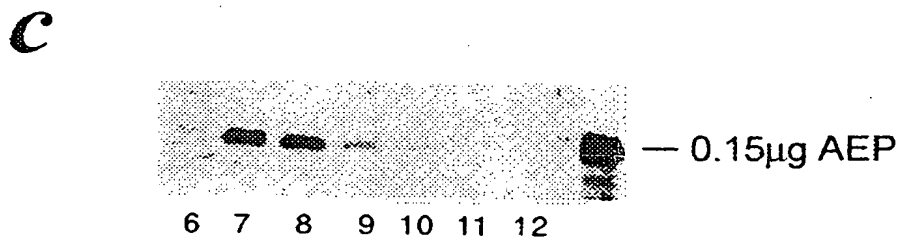
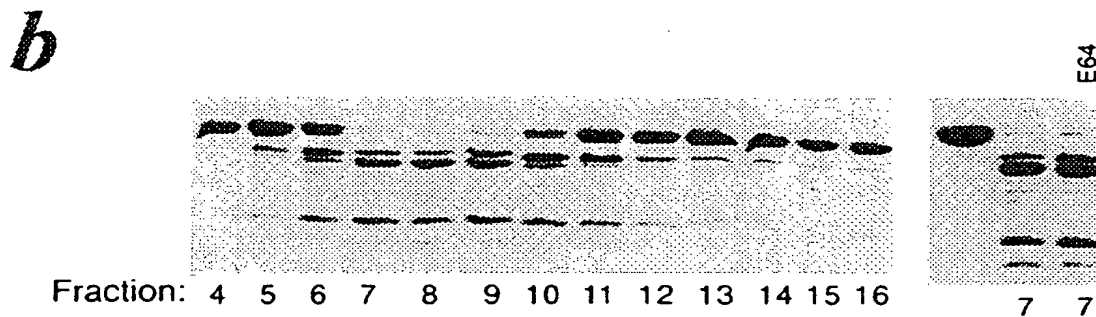
56. A composition comprising an inhibitor of asparaginyl endopeptidase and an inhibitor of cathepsin S.

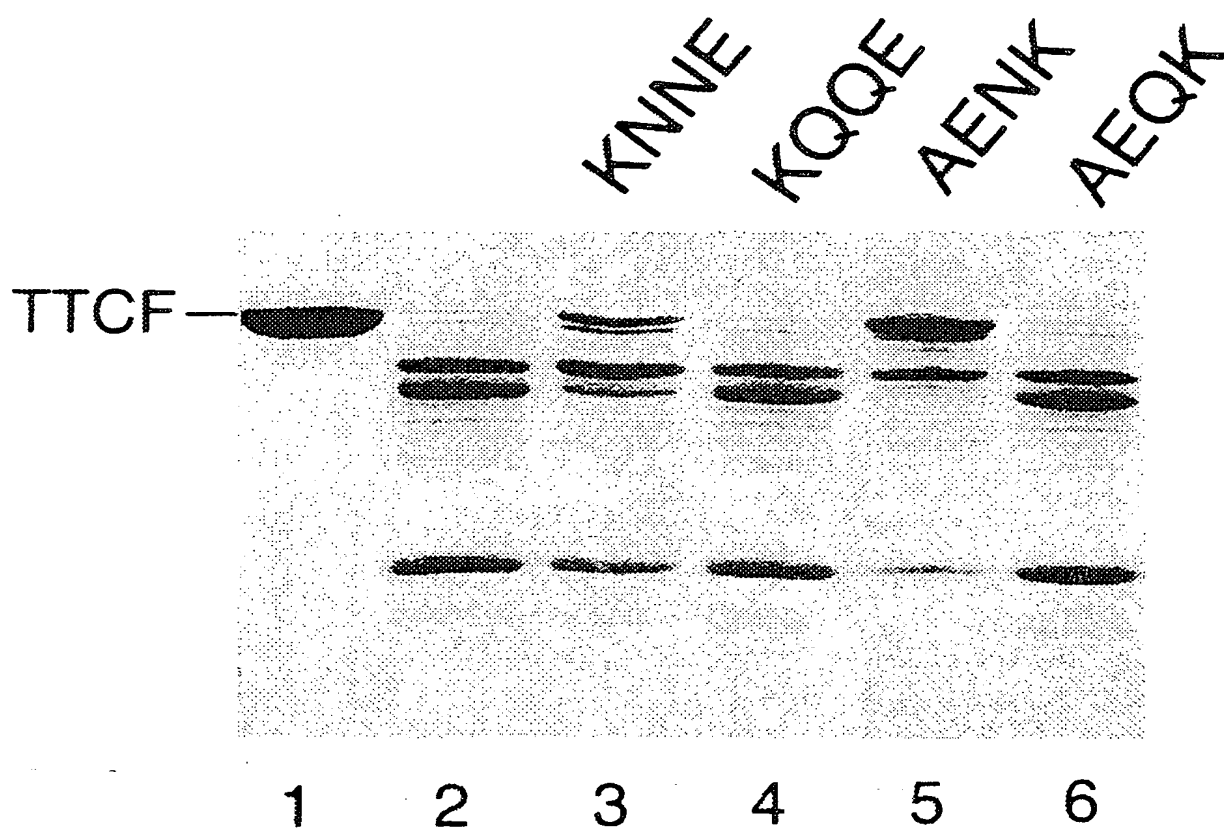
*Fig. 1 (page 1 of 2)*



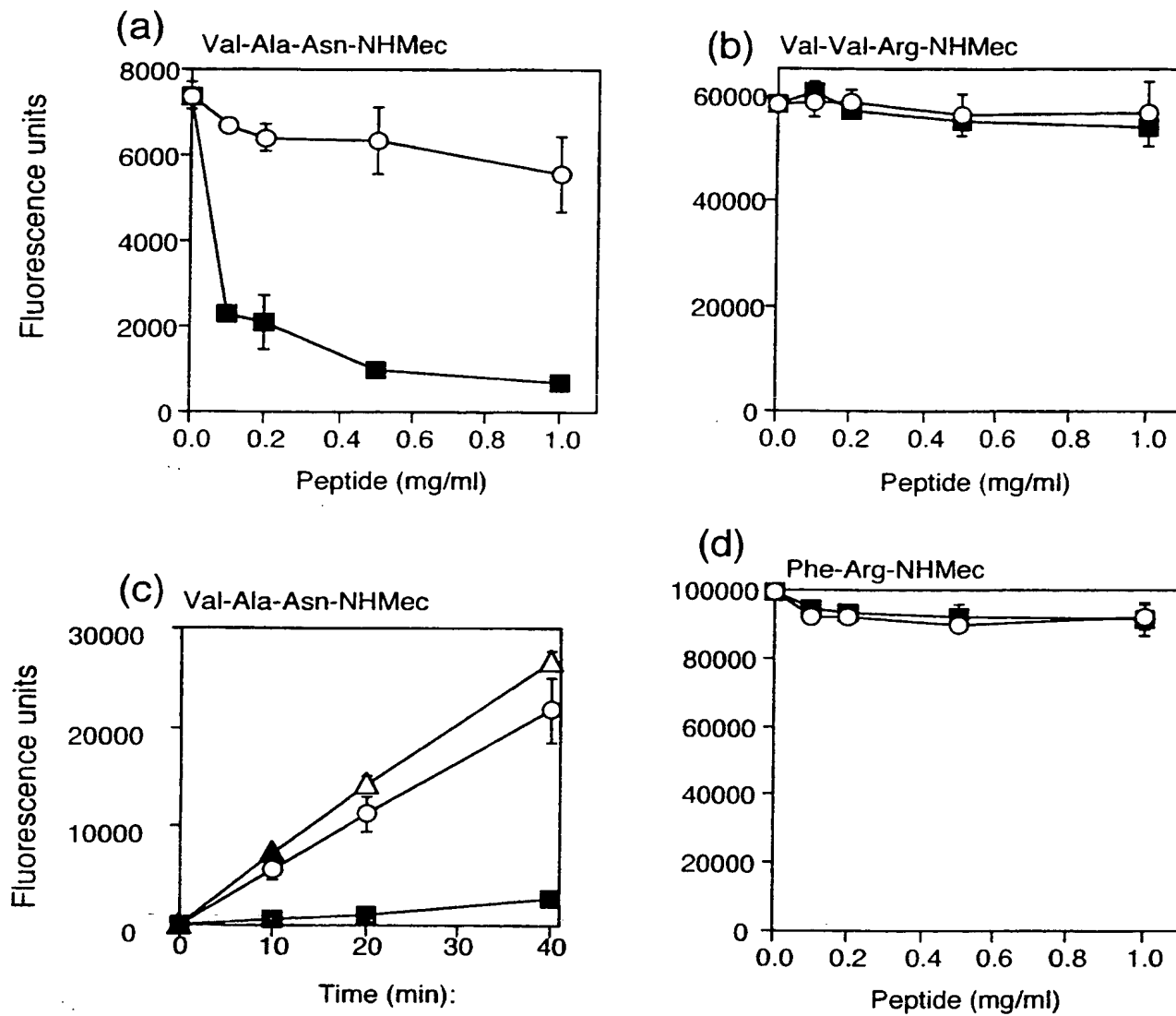
***Fig. 1 (page 2 of 2)***

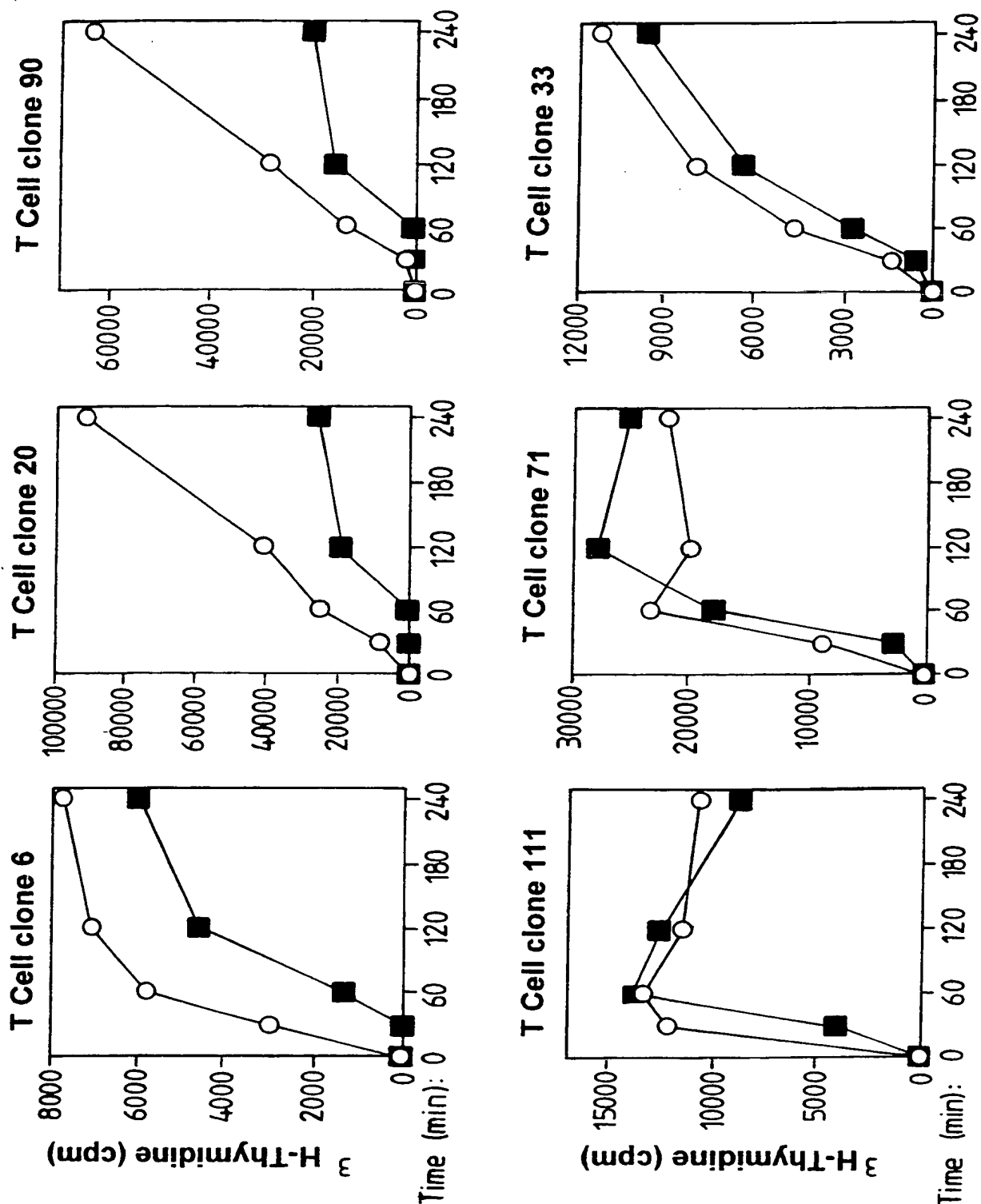


**Fig. 2**

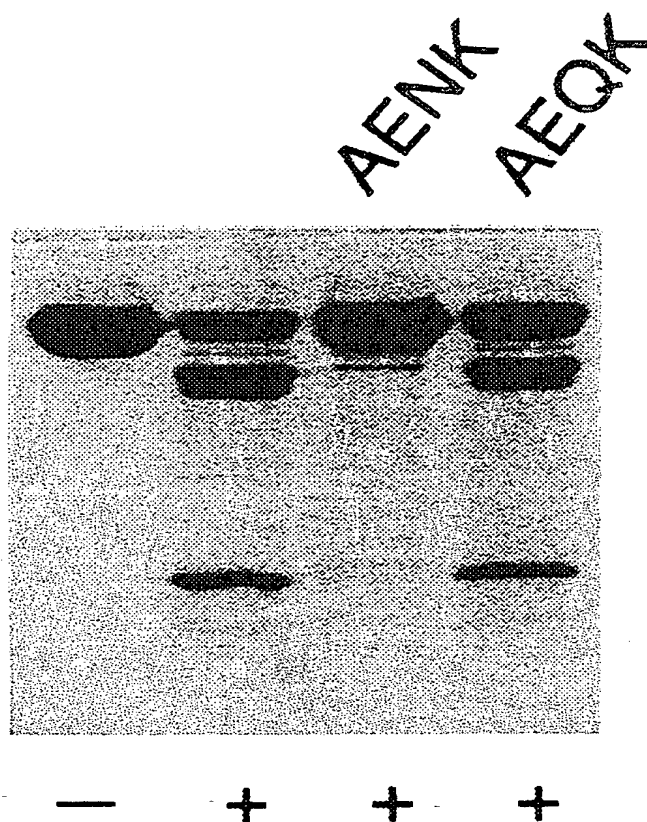


*Fig. 3a*

***Fig. 3b***

*Fig 3c*

6/24

*Fig. 3d*

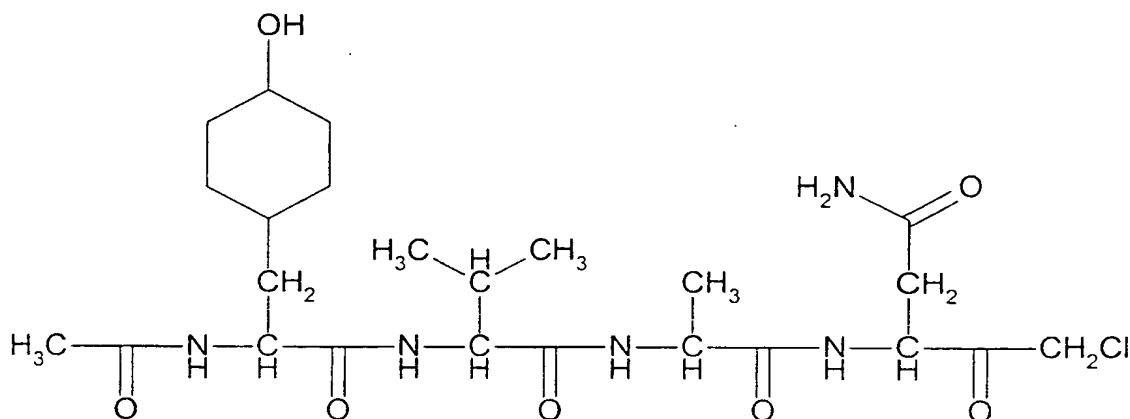
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Human B cell AEP  
 $\pm$  1.0 mg/ml peptide

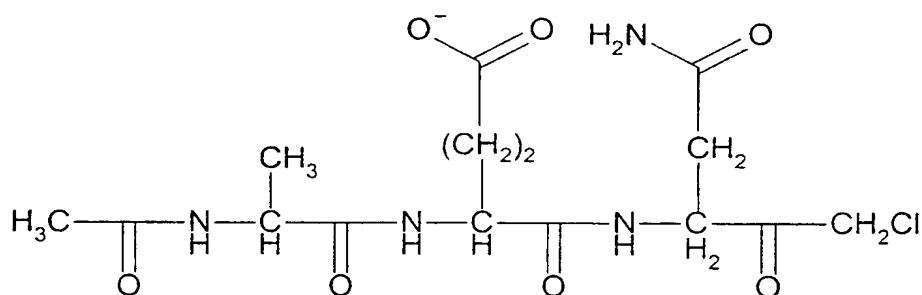
**Figure 4 (page 1 of 6)**

Peptidyl chloromethylketones (ref 5)

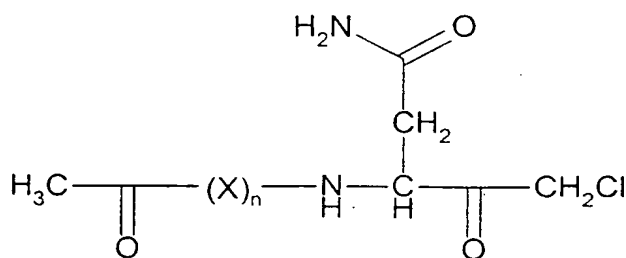
- (i) Acetyl-tyrosyl-valyl-alanyl-asparaginyl-chloromethylketone  
(analogous to ICE protease inhibitor YVAD-cmk)



- (ii) Acetyl-alanyl-glutamyl-asparaginyl-chloromethylketone



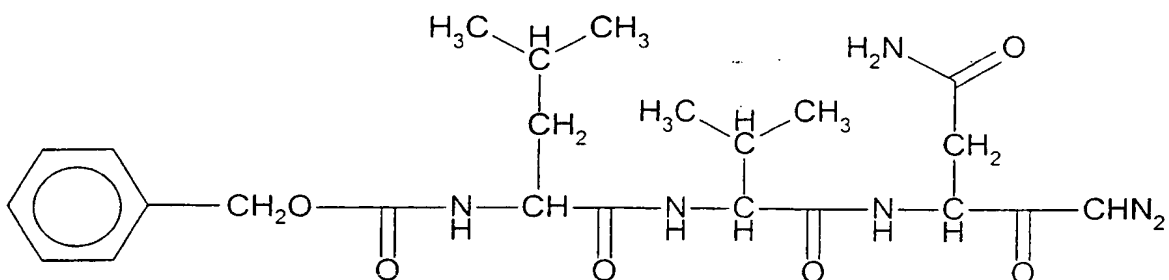
- (iii) Acetyl (or benzyloxycarbonyl)-(X)<sub>n</sub>-asparaginyl-chloromethylketone  
Where X = any amino acid



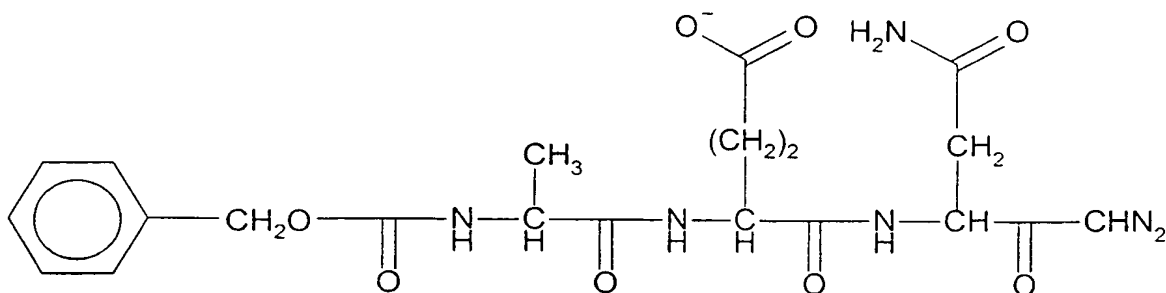
**Figure 4 (page 2 of 6)**

Peptidyl diazomethanes (ref 3,4)  
(have the general structure:  $R-C(=O)CHN_2$ )

(i) Benzyloxycarbonyl-leucyl-valyl-asparaginyl-diazomethane

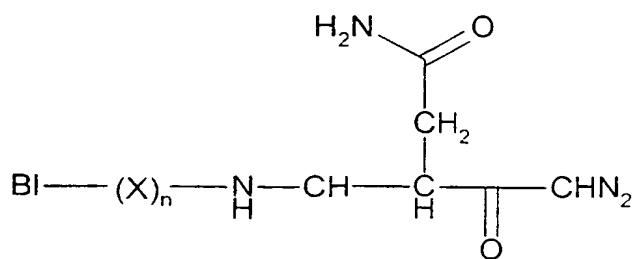


(ii) Alanyl-glutamyl-asparaginyl-diazomethane



(iii) Z-(X)<sub>n</sub>-asparaginyl-diazomethane

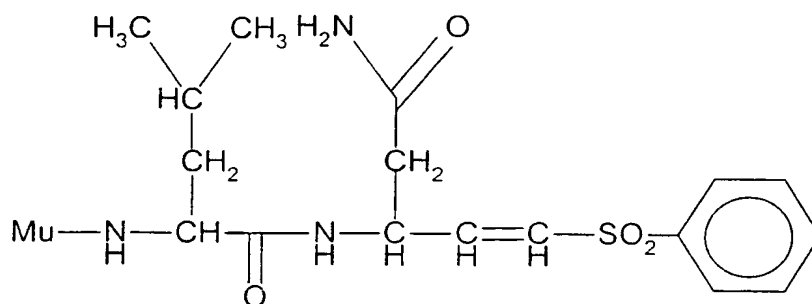
Where Bl = acetyl or benzyloxycarbonyl and X = any amino acid



**Figure 4 (page 3 of 6)**

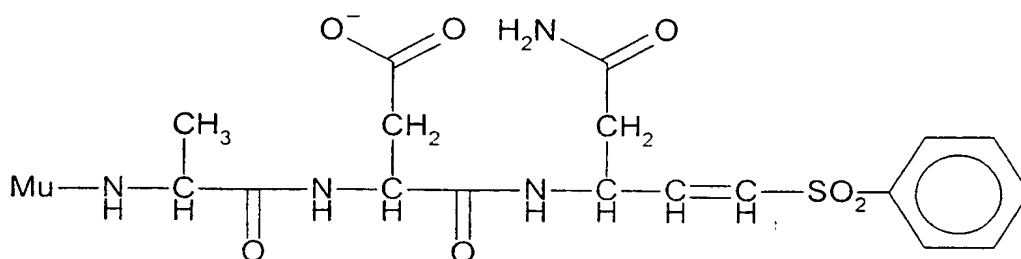
Peptidyl vinyl sulphones (ref 6)

- (i) Morpholinurea-leucyl-asparaginyl-vinylsulphone-phenyl  
(Acetyl or benzyloxycarbonyl can replace morpholinurea)



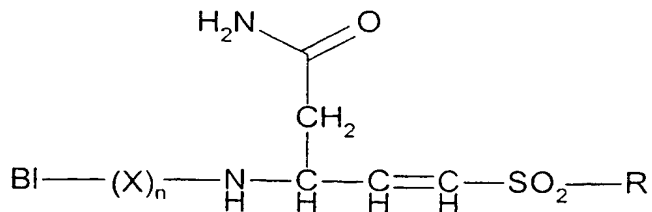
Mu - morpholinurea

- (ii) Morpholinurea-alanyl-glutamyl-asparaginyl-vinylsulphone-phenyl  
(Acetyl or benzyloxycarbonyl can replace morpholinurea)



- (iii) Bl-(X)<sub>n</sub>-asparaginyl-vinylsulphone-R

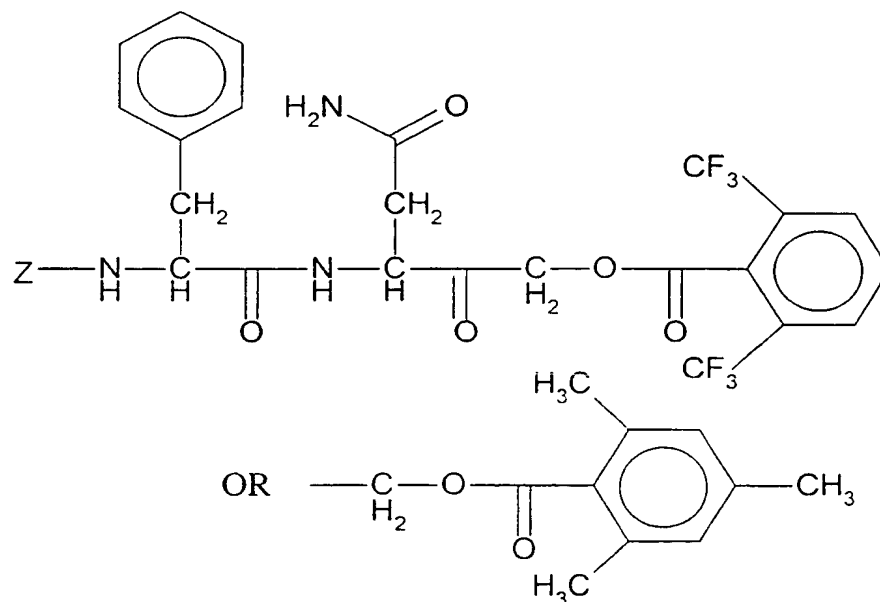
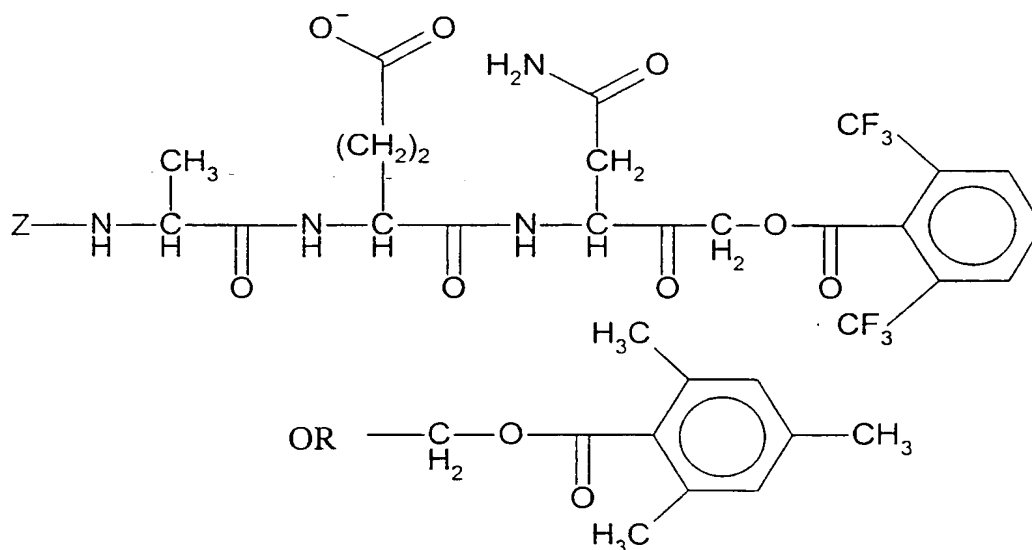
Where Bl = N-terminal blocking group (acetyl, morpholinurea or benzyloxycarbonyl), X = any amino acid  
and R = alkyl or aryl terminating group





**Figure 4 (page 4 of 6)**

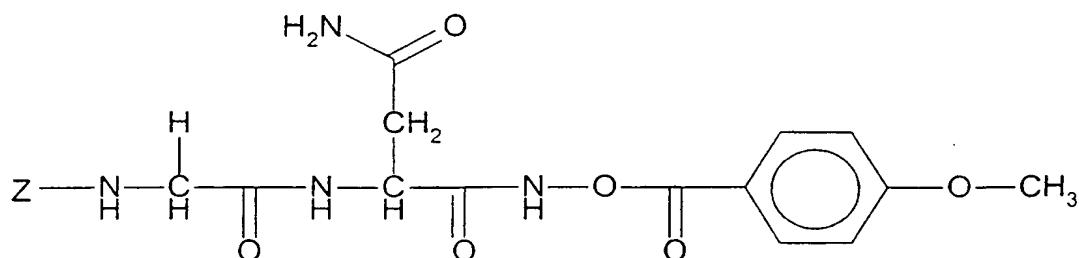
Peptidyl (acyloxy) methanes (ref 7)

(i) Z-Phenylalanyl-asparaginyl-CH<sub>2</sub>OCO-[2,6-(CF<sub>3</sub>)<sub>2</sub>Phenyl](ii) Z-Phenylalanyl-asparaginyl-CH<sub>2</sub>OCO-[2,4,6-(CH<sub>3</sub>)<sub>3</sub> Phenyl](iii) Z-alanyl-glutamyl-asparaginyl-CH<sub>2</sub>OCO-[2,6-(CF<sub>3</sub>)<sub>2</sub>Phenyl](iv) Z-alanyl-glutamyl-asparaginyl-CH<sub>2</sub>OCO-[2,4,6-(CH<sub>3</sub>)<sub>3</sub> Phenyl](v) Z-(X)<sub>n</sub>-asparaginyl-CH<sub>2</sub>OCO-RWhere X = any amino acid and R = [2,6-(CF<sub>3</sub>)<sub>2</sub>Phenyl] or [2,4,6-(CH<sub>3</sub>)<sub>3</sub> Phenyl] or other acyloxy methane group

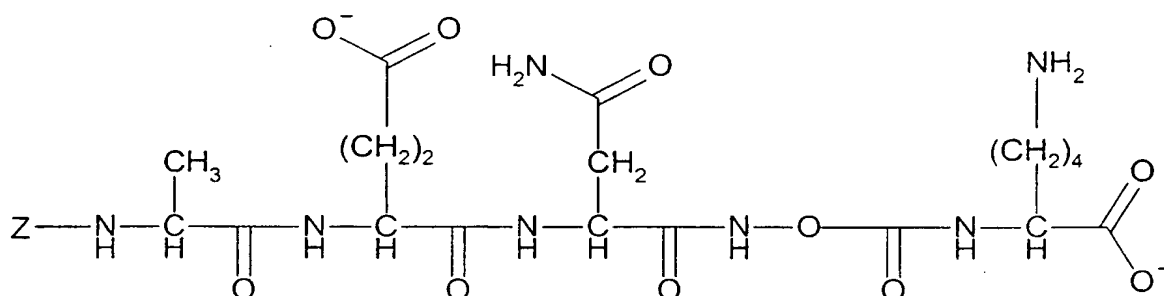
**Figure 4 (page 5 of 6)**

N,O-diacyl hydroxamates (ref 8)

(i) Z-Glycyl-asparaginyl-NHO-benzoyl(4-OCH<sub>3</sub>)

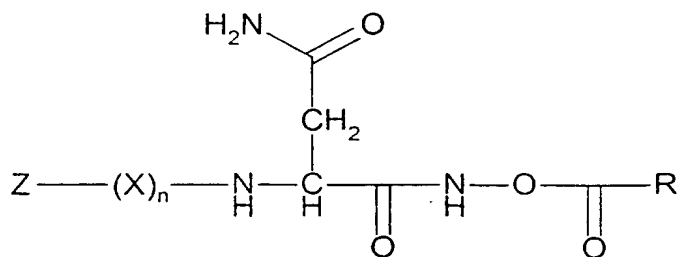


(ii) Z-alanyl-glutamyl-asparaginyl-NHO-CO-lysine-NH



(iv) Z-(X)<sub>n</sub>-asparaginyl-NHO-CO-R

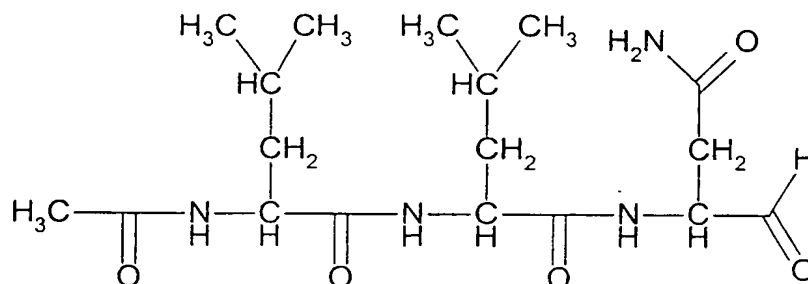
Where Z = benzyloxycarbonyl or other blocking group,  
X = any amino acid and R = any O-acyl group



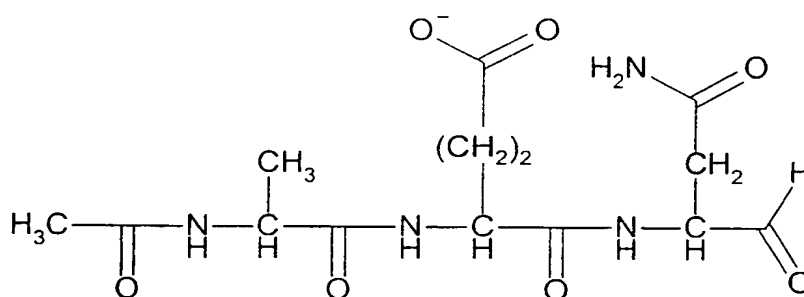
**Figure 4 (page 6 of 6)**

Peptide aldehydes (refs 1 &amp; 2)

(i) Acetyl-leucyl-leucyl-asparaginal



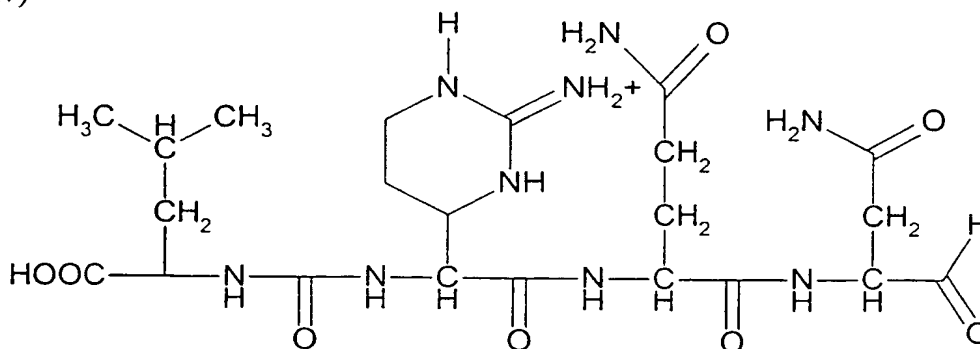
(ii) Acetyl-alanyl-glutamyl-asparaginal

(iv) Acetyl (or other blocking group)-(X)<sub>n</sub>-Asparaginal

-where X denotes any amino acid(s) in peptide linkage

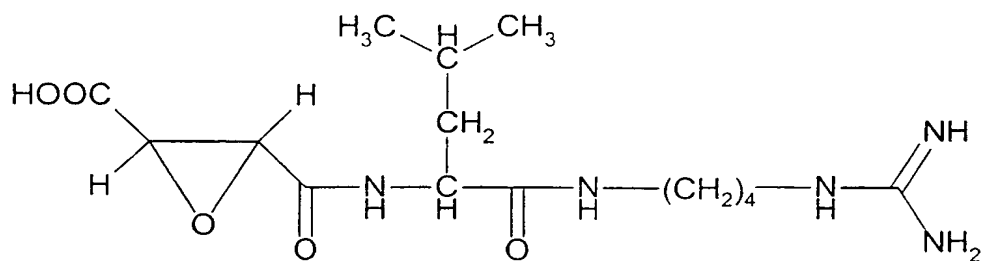
Elastinal also blocks AEP. A more specific variant would be:

(iv)



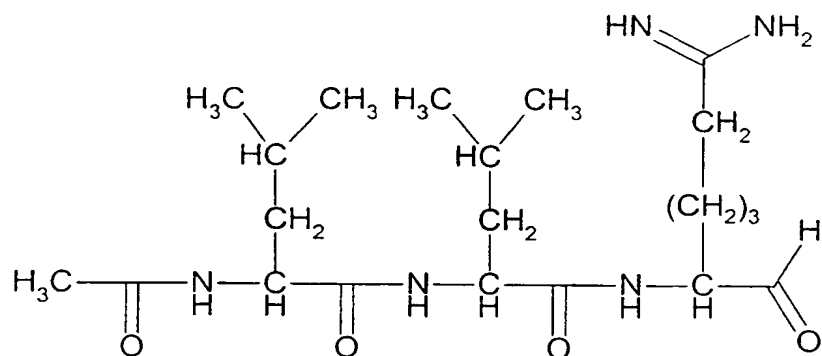
**Figure 5 (page 1 of 6)****E-64**

**Structure:** L-trans-epoxysuccinyl-leucylamide-(4-guanido)-butane or N-[N-(L-trans-carboxyoxiran-2-carbonyl)-L-leucyl]-agmatine



**Figure 5 (page 2 of 6)****Leupeptin**

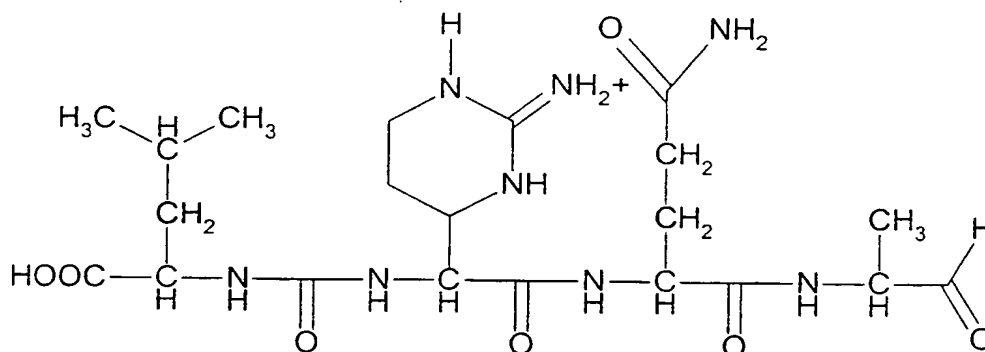
**Structure:** Acetyl-leucyl-leucyl arginal





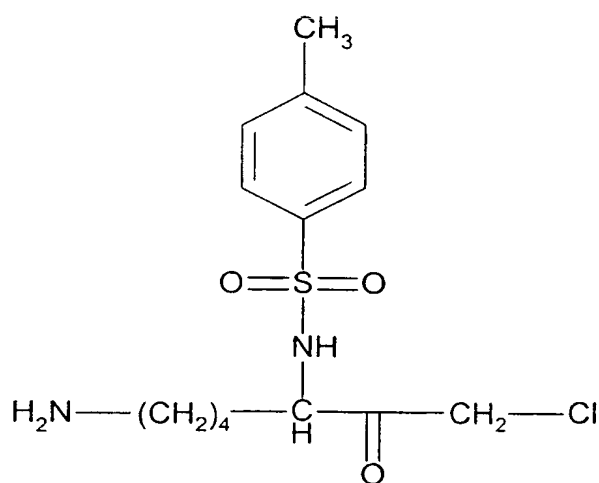
**Figure 5 (page 4 of 6)****Elastinal**

**Structure:** Leu-(Cap)-Gln-Ala-al,  
N-[(S)-1carboxy-isopentyl]-carbamoyl-alpha-(2-iminohexahydro-4(S)-  
pyrimidyl]-L-glycyl-L-glutamyl-L-alaninal



**Figure 5 (page 5 of 6)****TLCK**

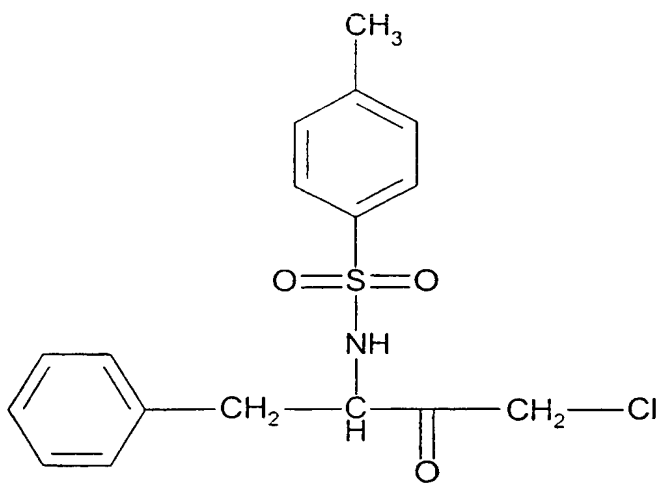
**Structure:** Tosyl Lysyl ChloromethylKetone:  
1-Chloro-3-tosylamido-7-amino-2-heptanone

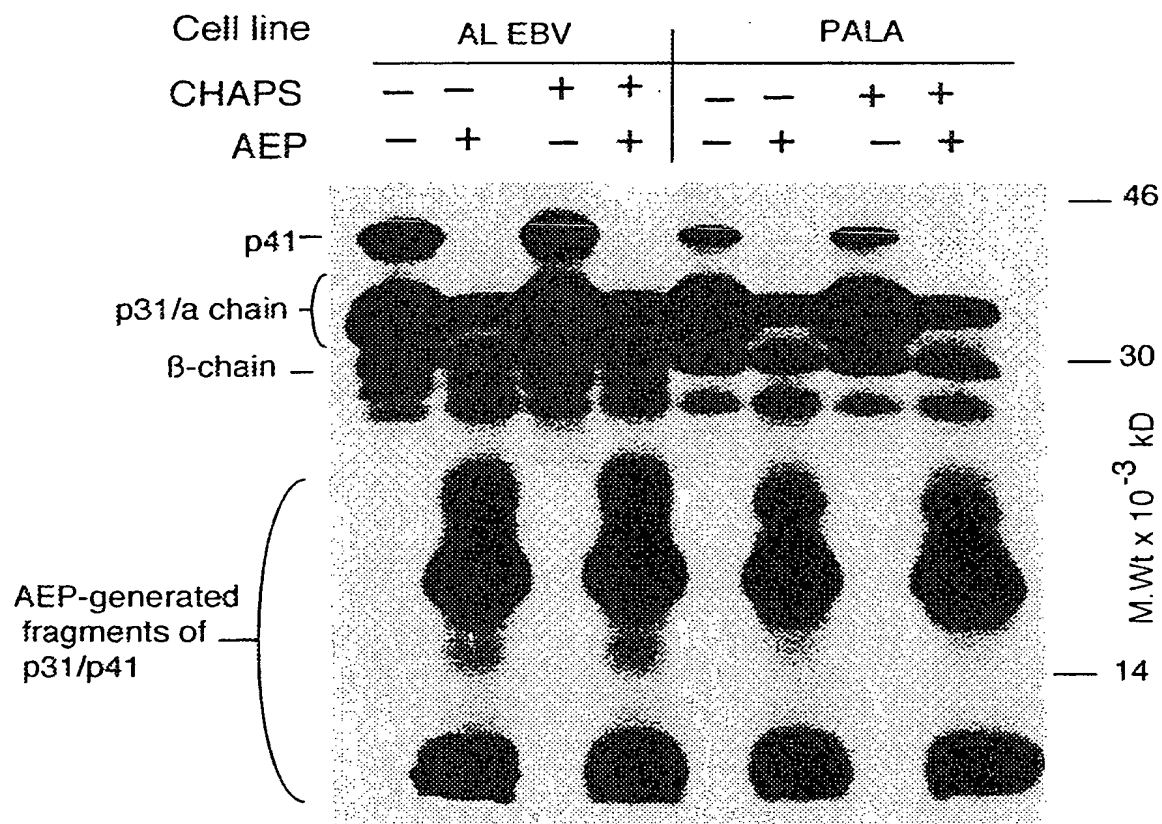




**Figure 5 (page 6 of 6)****TPCK**

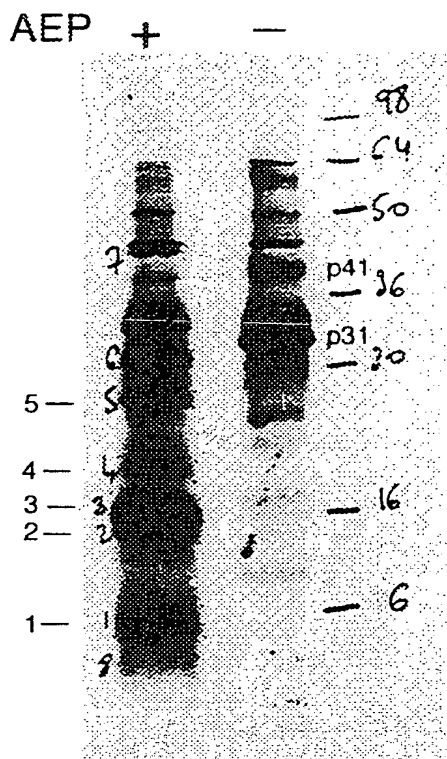
**Structure:** Tosyl Phenylalanyl ChloromethylKetone:  
1-Chloro-3-tosylamido-4-phenyl-2-butanone



*Fig. 6*

## AEP processing of the invariant chain

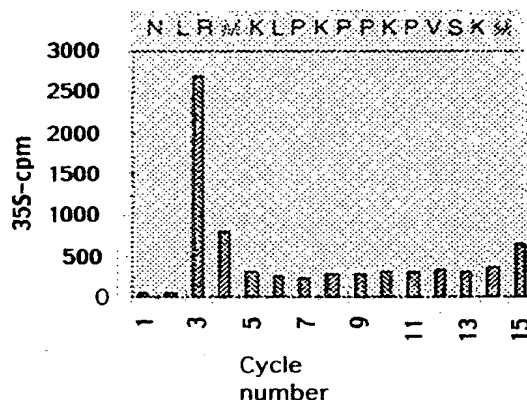
(a)

VIC  $\gamma$ 1 precipitate

(b) N-terminal sequence of bands below

- Band 1: N NEQLPM (Peak cycle 5/6)  
 Band 2: N TMETI (Peak cycle 2)  
 Band 3: N LRMK (Peak cycle 3)  
 Band 4: N LRMK  
 N NEQLPM (Peaks at cycles 3 & 6)  
 Band 5: N LRMK

Example raw data from band 5



(c)

MDDQ<sup>↓</sup>RD<sup>↓</sup>LISNNEQLPMLGR<sup>↓</sup>PGAPESKCSRGALYTGFSILVTLLLAGQATTAYF  
 QQQGR<sup>↓</sup>LDKLT<sup>↓</sup>VTSONLQLENLRMKLPKPPKPVSKMRMATPLL<sup>↓</sup>MOALPMGALPQG  
 QNATKYGNMTEDHVMHLLQNADPLKVYPPLKGSFPENLTHLKNTMETIDWKVFE  
 MHHWLLFEMSRHSLEQKPTDAPPKESLELEDPSGLGVTKQDLGPVPM

# Fig 8

## Identified AEP cleavage sites

### Tetanus toxin C fragment

RHIDN EEDID  
YTPNN EIDSF  
GNAFN NLDRI

### Ribonuclease

NGQTN CYQSY  
VACKN GQTNC

### Ovalbumin

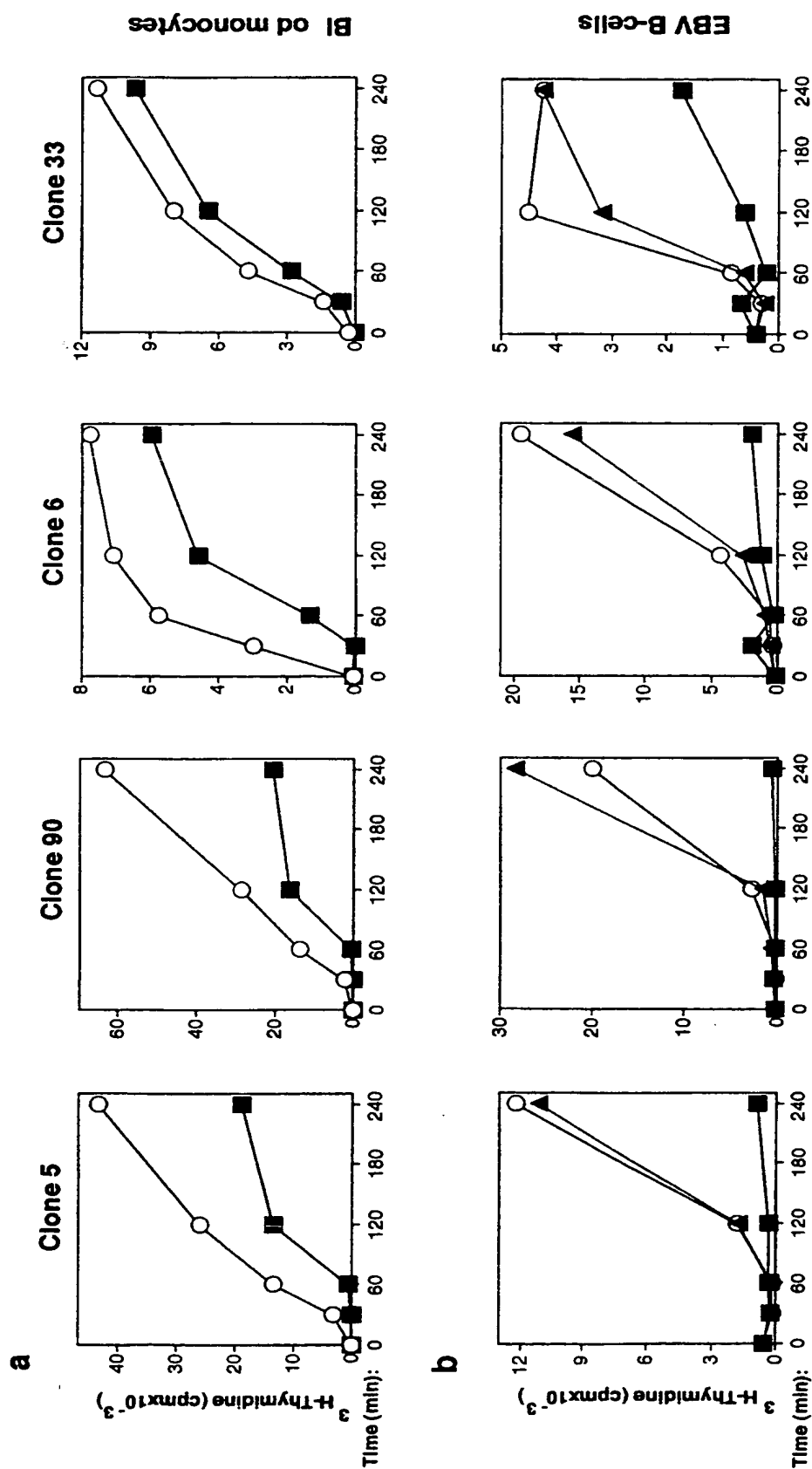
GTSVN VHSSL

### Hen Egg Lysozyme (preferred sites listed first)

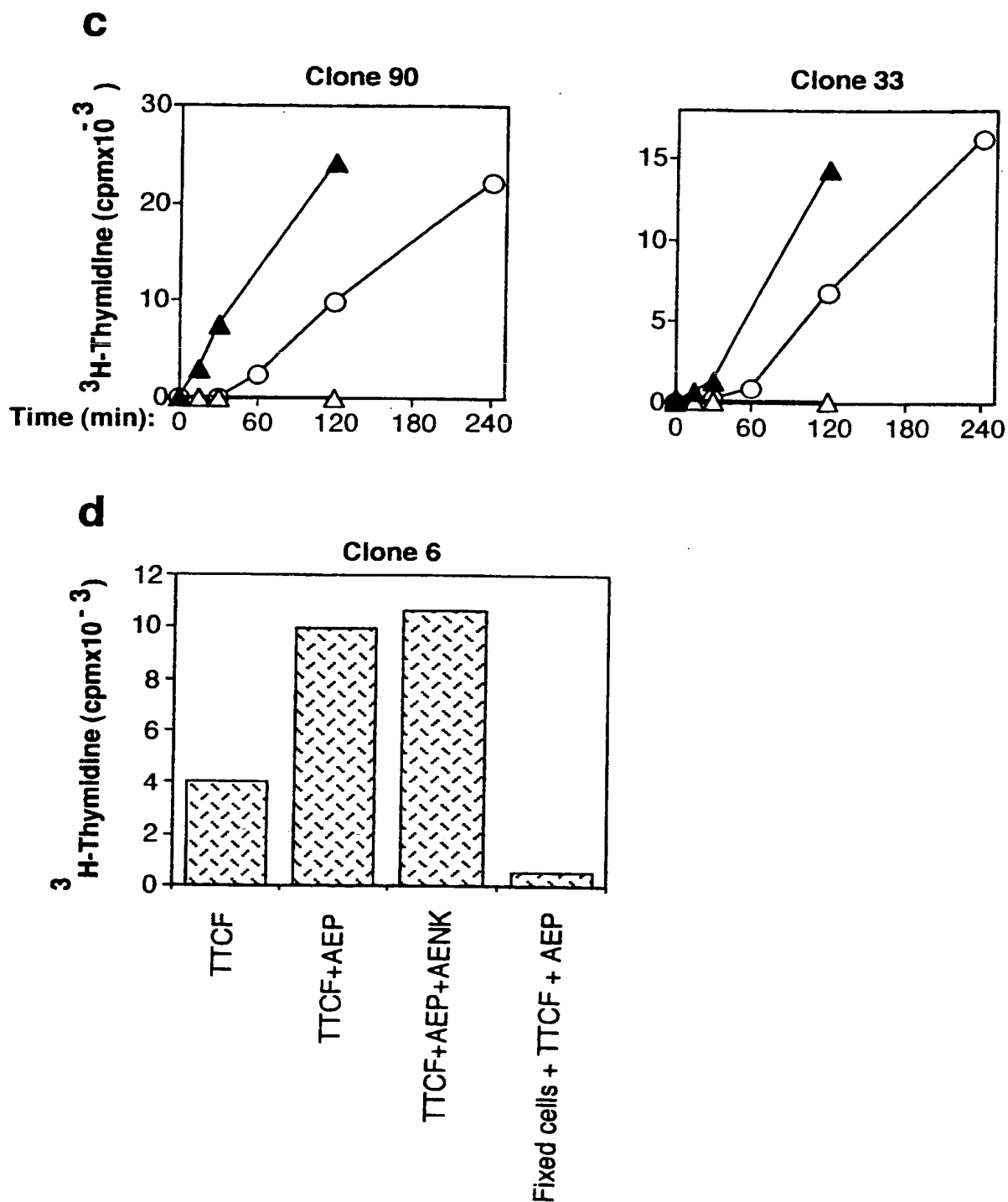
GNGMN AWVAW  
HGLDN YRGYS  
ILQIN SRWWC  
VSDGN GMNAW  
RWWCN DGRTP  
VAWRN RCKGT

### Transferrin peptide (622-642)

LFGSN VTDCS  
DCSGN FCLFR

*Fig. 9 (page 1 of 2)*

# *Fig. 9 (page 2 of 2)*



# INTERNATIONAL SEARCH REPORT

In International Application No

PCT/GB 99/00963

## A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C07K5/06 C07K5/08 A61K38/55 C07K14/705 G01N33/68  
A01K67/027

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C07K A61K G01N A01K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

| Category * | Citation of document, with indication, where appropriate, of the relevant passages   | Relevant to claim No. |
|------------|--|-----------------------|
| X          | CHEMICAL ABSTRACTS, vol. 128, no. 25,<br>22 June 1998 (1998-06-22)<br>Columbus, Ohio, US;<br>abstract no. 303708,<br>W C BENJAMIN ET AL.: "Caspase activation<br>in MCF7 cells responding to etoposide<br>treatment"<br>XP002113051<br>& MOL. PHARMACOL.,<br>vol. 53, no. 3, 1998, pages 446-450,<br>abstract<br><br>---<br>-/-- | 52,53                 |

☒ Further documents are listed in the continuation of box C.

☐ Patent family members are listed in annex.

### \* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

23 August 1999

Date of mailing of the international search report

03/09/1999

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2  
NL - 2280 HV Rijswijk  
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Fax: (+31-70) 340-3016

Authorized officer

Masturzo, P

## INTERNATIONAL SEARCH REPORT

International Application No

PCT/GB 99/00963

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

| Category * | Citation of document, with indication, where appropriate, of the relevant passages  | Relevant to claim No. |
|------------|---|-----------------------|
| X          | K K SHARMA ET AL.: "identification of new lens protease(s) using substrates having in vivo cleavage sites" BIOCHEMICAL AND BIOPHYSICAL RESEARCH COMMUNICATIONS., vol. 218, 1996, pages 365-370, XP002113049 ACADEMIC PRESS INC. ORLANDO, FL., US ISSN: 0006-291X the whole document<br>---  | 55                    |
| A          | J M CHEN ET AL.: "Cloning, isolation and characterization of mammalian legumain, an asparaginyl endopeptidase" JOURNAL OF BIOLOGICAL CHEMISTRY., vol. 272, no. 12, 21 March 1997 (1997-03-21), pages 80090-8098, XP002065429 AMERICAN SOCIETY OF BIOLOGICAL CHEMISTS, BALTIMORE, MD., US ISSN: 0021-9258 cited in the application the whole document<br>---         | 1-56                  |
| P,X        | B MANOURY T AL.: "An asparaginyl endopeptidase processes a microbial antigen for class II MHC presentation" NATURE., vol. 396, 17 December 1998 (1998-12-17), pages 695-699, XP002113050 MACMILLAN JOURNALS LTD. LONDON., GB ISSN: 0028-0836 the whole document<br>---  | 1-56                  |
| A          | CHEMICAL ABSTRACTS, vol. 119, no. 13, 27 September 1993 (1993-09-27) Columbus, Ohio, US; abstract no. 133992, A A KEMBHAVI ET AL.: "The two cysteine endopeptidases of legume seeds; purification and characterization by use of specific fluorometric assays" XP002113052 & ARCH. BIOCHEM. BIOPHYS., vol. 303, no. 2, 1993, pages 208-213, abstract<br>---<br>-/-- | 43-45                 |



## INTERNATIONAL SEARCH REPORT

International Application No

PCT/GB 99/00963

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

| Category * | Citation of document, with indication, where appropriate, of the relevant passages   | Relevant to claim No. |
|------------|--|-----------------------|
| X          | <p>CHEMICAL ABSTRACTS, vol. 121, no. 17,<br/>24 October 1994 (1994-10-24)<br/>Columbus, Ohio, US;<br/>abstract no. 202741,<br/>M J ANDROLEWICZ ET AL.: "Human<br/>transporters associated with antigen<br/>processing possess a promiscuous binding<br/>site"<br/>XP002113053<br/>&amp; IMMUNITY,<br/>vol. 1, no. 1, 1994, pages 7-14,<br/>abstract</p> <p>-----</p> | 55                    |

# INTERNATIONAL SEARCH REPORT

International application No.

PCT/GB 99/ 00963

## Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 1-21  
because they relate to subject matter not required to be searched by this Authority, namely:  
Remark: Although claims 1-21 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. ☐ Claims Nos.:  
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

## Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

### Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.